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(54) Title: METHODS OF DIAGNOSIS OF LUNG CANCER, COMPOSITIONS AND METHODS OF SCREENING FOR MOD-**ULATORS OF LUNG CANCER**

METHODS OF DIAGNOSIS OF LUNG CANCER, COMPOSITIONS AND METHODS OF SCREENING FOR MODULATORS OF LUNG CANCER

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CROSS-REFERENCES TO RELATED APPLICATIONS

This application is related to USSN 60/284,770, filed April 18, 2001; USSN 60/290,492, filed May 10, 2001; USSN 60/334,370, filed November 29, 2001; USSN 60/339,245, filed November 9, 2001; USSN 60/350,666, filed November 13, 2001; and USSN 60/xxx,xxx, filed April 12, 2002 (Docket OMNI-002P); each of which is incorporated herein by reference in its entirety.

FIELD OF THE INVENTION

The invention relates to the identification of nucleic acid and protein expression profiles and nucleic acids, products, and antibodies thereto that are involved in lung cancer; and to the use of such expression profiles and compositions in diagnosis and therapy of lung cancer. The invention further relates to methods for identifying and using agents and/or targets that inhibit lung cancer or related conditions.

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BACKGROUND OF THE INVENTION

Lung cancer is the second most commonly occurring cancer in the United States and is the leading cause of cancer-related death. It is estimated that there are over 160,000 new cases of lung cancer in the United States every year. Of those who are diagnosed with lung cancer, 86 percent will die within five years. Lung cancer is the most common visceral cancer in men and accounts for nearly one third of all cancer deaths in both men and women. In fact, lung cancer accounts for 7% of all deaths, due to any cause, in both men and women.

Smoking is the primary cause of lung cancer, with more than 80% of lung cancers resulting from smoking. About 400 to 500 separate gaseous substances are present in the smoke of a non-filter cigarette. The most noteworthy substances include nitrogen oxides, hydrogen cyanide, formaldehyde, benzene, and toluene. The particles present in cigarette smoke contain at least 3,500 individual compounds such as nicotine, tobacco alkaloids (nornicotine, anatabine, anabasine), polycyclic aromatic hydrocarbons (e.g., benzo(a)pyrene, B(a)P), naphthalenes, aromatic amines, phenols, and tobacco-specific nitrosamines.

Tobacco-specific nitrosamines are formed during tobacco curing and processing, and are suspected of causing lung cancer in humans. In rodent studies, regardless of the where or how it is applied, the tobacco-specific nitrosamine known as NNK produces lung adenomas and lung adenocarcinomas. The tobacco-specific nitrosamine known as NNAL also produces lung adenocarcinomas in rodents.

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Many of the chemicals found in cigarette smoke also affect the nonsmoker inhaling "secondhand" or sidestream smoke. Indeed, the smoke inhaled by non-smokers has a chemical composition similar to the smoke inhaled by smokers, but, importantly, the concentrations of the carcinogenic tobacco-specific nitrosamines are present in higher concentrations in second hand smoke. For this and other reasons, "passive smoking" is an important cause of lung cancer, causing as many as 3,000 lung cancer deaths in nonsmokers each year.

In addition to smoking, other factors thought to be causes of lung cancer include onthe-job exposure to carcinogens such as asbestos and uranium, exposure to chemical hazards such as radon, polycyclic aromatic hydrocarbons, chromium, nickel, and inorganic arsenic, genetic factors, and diet.

Histological classification of various lung cancers define the types of cancer that begin in the lung. See, e.g., Travis, et al. (1999) <u>Histological Typing of Lung and Pleural Tumours</u> (International Histological Classification of Tumours, No 1. Four major cell types make up more than 88% of all primary lung neoplasms. These are: squamous or epidermoid carcinoma, small cell (also called oat cell) carcinoma, adenocarcinoma, and large cell (also called large cell anaplastic) carcinoma. The remainder include undifferentiated carcinomas, carcinoids, bronchial gland tumors, and other rarer types. The various cell types have different natural histories and responses to therapy, and, thus, a correct histologic diagnosis is the first step of effective treatment.

Small cell lung cancer (SCLC) accounts for 18-25% of all lung cancers, and occurs less frequently than non-small cell lung cancers, and generally spread to distant organs more rapidly than non-small cell lung cancer. In general, at the time of presentation small cell lung cancers have already spread beyond the beyond the bounds where surgery and curative intent can be undertaken. Hoever, if identified early enough, these cancers are often responsive to chemotherapy and thoracic radiation treatment.

Non-small cell lung cancers (NSCLC) are the more frequently occurring form of lung cancer. They comprise squamous cell carcinoma, adenocarcinoma, and large cell carcinoma

and account for more than 75% of all lung cancers. Non-small cell tumors that are localized at the time of presentation can sometimes be cured with surgery and/or radiotherapy, but usually are not identified until significant metastasis has occurred, which are typically not very responsive to surgical, chemotherapy, or radiation treatment..

The screening of asymptomatic persons at high risk for lung cancer has often proven ineffective. In general, only 5 to 15 percent of lung cancer patients have their disease detected while they are asymptomatic. Of course, early detection and treatment are critical factors in the fight against lung cancer. The average survival rate is 49% for those whose cancer is detected early, before the cancer has spread from the lung. Lung cancer often spreads outside of the lung, and it may have spread to the bones or brain by the time it is diagnosed. While the prognosis may be better for lung cancers that are detected early, because of the lack of effective curative treatments, early detection does not necessarily alter the total death rate from lung cancer.

Thus, methods for diagnosis and prognosis of lung cancer and effective treatment of lung cancer would be desirable. Accordingly, provided herein are methods that can be used in diagnosis and prognosis of lung cancer. Further provided are methods that can be used to screen candidate therapeutic agents for the ability to modulate, e.g., treat, lung cancer. Additionally, provided herein are molecular targets and compositions for therapeutic intervention in lung disease and other metastatic cancers.

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SUMMARY OF THE INVENTION

The present invention provides nucleotide sequences of genes that are up- and down-regulated in lung cancer cells. Such genes are useful for diagnostic purposes, and also as targets for screening for therapeutic compounds that modulate lung cancer, such as antibodies. The methods of detecting nucleic acids of the invention or their encoded proteins can be used for a number of purposes. Examples include early detection of lung cancers, monitoring and early detection of relapse following treatment of lung cancers, monitoring response to therapy of lung cancers, determining prognosis of lung cancers, directing therapy of lung cancers, selecting patients for postoperative chemotherapy or radiation therapy, selecting therapy, determining tumor prognosis, treatment, or response to treatment, and early detection of precancerous lesions of the lung. Examples of benign or precancerous lesions include: atelectasis, emphysema, brochitis, chronic obstructive pulmonary disease, fibrosis, hypersensitivity pneumonitis (HP), interstitial pulmonary fibrosis (IPF), asthma, and

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bronchiectasis. Other aspects of the invention will become apparent to the skilled artisan by
the following description of the invention.

In one aspect, the present invention provides a method of detecting a lung cancer-associated transcript in a cell from a patient, the method comprising contacting a biological sample from the patient with a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16. Alternatively, the sample may be contacted with a specific binding reagent, e.g., antibody.

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In one embodiment, the polynucleotide selectively hybridizes to a sequence at least 95% identical to a sequence as shown in Tables 1A-16. In another embodiment, the polynucleotide comprises a sequence as shown in Tables 1A-16.

In one embodiment, the biological sample is a tissue sample, or a body fluid. In another embodiment, the biological sample comprises isolated nucleic acids, e.g., mRNA.

In one embodiment, the polynucleotide is labeled, e.g., with a fluorescent label. In one embodiment, the polynucleotide is immobilized on a solid surface. In one embodiment, the patient is undergoing a therapeutic regimen to treat lung cancer. In another embodiment, the patient is suspected of having lung cancer. In one embodiment, the patient is a primate, e.g., a human.

In one embodiment, the method further comprises the step of amplifying nucleic acids before the step of contacting the biological sample with the polynucleotide.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated transcript in the biological sample by contacting the biological sample with a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, thereby monitoring the efficacy of the therapy. Or the sample may be evaluated for protein, e.g., contacting the sample with an antibody.

In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated transcript to a level of the lung cancer-associated transcript in a biological sample from the patient prior to, or earlier in, the therapeutic treatment. Or the sample may be evaluated for comparison of protein.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a

biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated antibody in the biological sample by contacting the biological sample with a polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, wherein the polypeptide specifically binds to the lung cancer-associated antibody, thereby monitoring the efficacy of the therapy.

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In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated antibody to a level of the lung cancer-associated antibody in a biological sample from the patient prior to, or earlier in, the therapeutic treatment.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated polypeptide in the biological sample by contacting the biological sample with an antibody, wherein the antibody specifically binds to a polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, thereby monitoring the efficacy of the therapy.

In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated polypeptide to a level of the lung cancer-associated polypeptide in a biological sample from the patient prior to, or earlier in, the therapeutic treatment. In one aspect, the present invention provides an isolated nucleic acid molecule consisting of a polynucleotide sequence as shown in Tables 1A-16. In one embodiment, an expression vector or cell comprises the isolated nucleic acid. In one aspect, the present invention provides an isolated polypeptide which is encoded by a nucleic acid molecule having polynucleotide sequence as shown in Tables 1A-16.

In another aspect, the present invention provides an antibody that specifically binds to an isolated polypeptide which is encoded by a nucleic acid molecule having polynucleotide sequence as shown in Tables 1A-16. In one embodiment, the antibody is conjugated to an effector component, e.g., a fluorescent label, a radioisotope or a cytotoxic chemical. In one embodiment, the antibody is an antibody fragment. In another embodiment, the antibody is humanized.

In one aspect, the present invention provides a method of detecting lung cancer in a a patient, the method comprising contacting a biological sample from the patient with an antibody or protein as described herein.

In another aspect, the present invention provides a method of detecting antibodies specific to a lung cancer gene in a patient, the method comprising contacting a biological sample from the patient with a polypeptide encoded by a nucleic acid comprises a sequence from Tables 1A-16.

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In another aspect, the present invention provides a method for identifying a compound that modulates a lung cancer-associated polypeptide, the method comprising the steps of: (i) contacting the compound with a lung cancer-associated polypeptide, the polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16; and (ii) determining the functional effect of the compound upon the polypeptide.

In one embodiment, the functional effect is a physical effect, an enzymatic effect, or a chemical effect. In one embodiment, the polypeptide is expressed in a eukaryotic host cell or cell membrane. In another embodiment, the polypeptide is recombinant. In one embodiment, the functional effect is determined by measuring ligand binding to the polypeptide.

In another aspect, the present invention provides a method of inhibiting proliferation or another critical process of a lung cancer-associated cell to treat lung cancer in a patient, the method comprising the step of administering to the subject a therapeutically effective amount of a compound identified as described herein. In one embodiment, the compound is an antibody.

In another aspect, the present invention provides a drug screening assay comprising the steps of: (i) administering a test compound to a mammal having lung cancer or a cell isolated therefrom; (ii) comparing the level of gene expression of a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16 in a treated cell or mammal with the level of gene expression of the polynucleotide in a control cell or mammal, wherein a test compound that modulates the level of expression of the polynucleotide is a candidate for the treatment of lung cancer.

In one embodiment, the control is a mammal with lung cancer or a cell therefrom that has not been treated with the test compound. In another embodiment, the control is a normal cell or mammal, or a non-malignant lung disease.

In another aspect, the present invention provides a method for treating a mammal having lung cancer comprising administering a compound identified by the assay described herein.

In another aspect, the present invention provides a pharmaceutical composition for treating a mammal having lung cancer, the composition comprising a compound identified by the assay described herein and a physiologically acceptable excipient.

DETAILED DESCRIPTION OF THE INVENTION

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In accordance with the objects outlined above, the present invention provides novel methods for diagnosis and treatment of lung disease or cancer, as well as methods for screening for compositions which modulate lung cancer. "Treatment, monitoring, detection or modulation of lung disease or cancer" includes treatment, monitoring, detection, or modulation of lung disease in those patients who have lung disease (whether malignant or non-malignant, e.g., emphysema, bronchitis, or fibrosis) as well as patients with lung cancers in which gene expression from a gene in Tables 1A-16 is increased or decreased, indicating that the subject is more likely to have disease. In particular, while these targets are identified primarily from lung cancer samples, these same targets are likely to be similarly found in analyses of other medical conditions. These other conditions may result from similar pathological processes which affect similar tissues, e.g., lung cancer, small cell lung carcinoma (oat cell carcinoma), non-small cell carcinomas (e.g., squamous cell carcinoma, adenocarcinoma, large cell lung carcinoma, carcinoid, granulomatous), fibrosis (idiopathic pulmonary fibrosis (IPF), hypersensitivity pneumonitis (HP), interstitial pneumonitis, nonspecific idiopathic pneumonitis (NSIP)), chronic obstructive pulmonary disease (COPD, e.g., emphysema, chronic bronchitis), asthma, bronchiectasis, and esophageal cancer. See, e.g., the NCI webpage and USSN 60/347,349 and USSN 60/xxx,xxx (docket LFBR-001-1P, filed March 29, 2002), each of which is incorporated herein by reference. The treatment may be of lung cancer or related condition itself, or treatment of metastasis.

In particular, identification of markers selectively expressed on these cancers allows for use of that expression in diagnostic, prognostic, or therapeutic methods. As such, the invention defines various compositions, e.g., nucleic acids, polypeptides, antibodies, and small molecule agonists/antagonists, which will be useful to selectively identify those markers. For example, therapeutic methods may take the form of protein therapeutics which use the marker expression for selective localization or modulation of function (for those markers which have a causative disease effect), for vaccines, identification of binding partners, or antagonism, e.g., using antisense or RNAi. The markers may be useful for molecular characterization of subsets of lung diseases, which subsets may actually require

very different treatments. Moreover, the markers may also be important in related diseases to the specific cancers, e.g., which affect similar tissues in non-malignant diseases, or have similar mechanisms of induction/maintenance. Metastatic processes or characteristics may also be targeted. Diagnostic and prognostic uses are made available, e.g., to subset related but distinct diseases, or to determine treatment strategy. The detection methods may be based upon nucleic acid, e.g., PCR or hybridization techniques, or protein, e.g., ELISA, imaging, IHC, etc. The diagnosis may be qualitative or quantitative, and may detect increases or decreases in expression levels.

Tables 1A-16 provide unigene cluster identification numbers for the nucleotide sequence of genes that exhibit increased or decreased expression in lung cancer samples. The tables also provide an exemplar accession number that provides a nucleotide sequence that is part of the unigene cluster. In Table 1A, genes marked as "target 1" or "target 2" are particularly useful as therapeutic targets. Genes marked as "target 3" are particularly useful as diagnostic markers. Genes marked as "chron" are upregulated in chronically diseased lung (e.g., emphysema, bronchitis, fibrosis) relative to lung tumors and normal tissue. In certain analyses, the ratio for the "chron" category was determined using the 70th percentile of chronically diseases lung samples divided by the 90th percentile of lung tumor samples divided by the 90th percentile of lung tumor samples divided by the 90th percentile of normal lung samples.

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Definitions

The term "lung cancer protein" or "lung cancer polynucleotide" or "lung cancer-associated transcript" refers to nucleic acid and polypeptide polymorphic variants, alleles, mutants, and interspecies homologs that: (1) have a nucleotide sequence that has greater than about 60% nucleotide sequence identity, 65%, 70%, 75%, 80%, 85%, 90%, preferably 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% or greater nucleotide sequence identity, preferably over a region of over a region of at least about 25, 50, 100, 200, 500, 1000, or more nucleotides, to a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16; (2) bind to antibodies, e.g., polyclonal antibodies, raised against an immunogen comprising an amino acid sequence encoded by a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16, and conservatively modified variants thereof; (3) specifically hybridize under stringent hybridization conditions to a nucleic acid sequence, or the complement thereof of Tables 1A-16 and conservatively modified variants thereof; or (4)

have an amino acid sequence that has greater than about 60% amino acid sequence identity, 65%, 70%, 75%, 80%, 85%, 90%, preferably 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% or greater amino sequence identity, preferably over a region of over a region of at least about 25, 50, 100, 200, 500, 1000, or more amino acid, to an amino acid sequence encoded by a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16. A polynucleotide or polypeptide sequence is typically from a mammal including, but not limited to, primate, e.g., human; rodent, e.g., rat, mouse, hamster; cow, pig, horse, sheep, or other mammal. A "lung cancer polypeptide" and a "lung cancer polynucleotide," include both naturally occurring or recombinant forms.

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A "full length" lung cancer protein or nucleic acid refers to a lung cancer polypeptide or polynucleotide sequence, or a variant thereof, that contains the elements normally contained in one or more naturally occurring, wild type lung cancer polynucleotide or polypeptide sequences. The "full length" may be prior to, or after, various stages of post-translational processing or splicing, including alternative splicing.

"Biological sample" as used herein is a sample of biological tissue or fluid that contains nucleic acids or polypeptides, e.g., of a lung cancer protein, polynucleotide, or transcript. Such samples include, but are not limited to, tissue isolated from primates, e.g., humans, or rodents, e.g., mice, and rats. Biological samples may also include sections of tissues such as biopsy and autopsy samples, frozen sections taken for histologic purposes, archival materials, blood, plasma, serum, sputum, stool, tears, mucus, hair, skin, etc. Biological samples also include explants and primary and/or transformed cell cultures derived from patient tissues. A biological sample is typically obtained from a eukaryotic organism, most preferably a mammal such as a primate, e.g., chimpanzee or human; cow; dog; cat; a rodent, e.g., guinea pig, rat, mouse; rabbit; or other mammal; or a bird; reptile; fish. Livestock and domestic animals are of interest.

"Providing a biological sample" means to obtain a biological sample for use in methods described in this invention. Most often, this will be done by removing a sample of cells from an animal, but can also be accomplished by using previously isolated cells (e.g., isolated by another person, at another time, and/or for another purpose), or by performing the methods of the invention in vivo. Archival tissues or materials, having treatment or outcome history, will be particularly useful.

The terms "identical" or percent "identity," in the context of two or more nucleic acids or polypeptide sequences, refer to two or more sequences or subsequences that are the

same or have a specified percentage of amino acid residues or nucleotides that are the same (e.g., about 60% identity, preferably 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or higher identity over a specified region, when compared and aligned for maximum correspondence over a comparison window or designated region) as measured using, e.g., a BLAST or BLAST 2.0 sequence comparison algorithms with default parameters described below, or by manual alignment and visual inspection (see, e.g., NCBI web site http://www.ncbi.nlm.nih.gov/BLAST/ or the like). Such sequences are then said to be "substantially identical." This definition also refers to, or may be applied to, the complement of a test sequence. The definition also includes sequences that have deletions and/or insertions, substitutions, and naturally occurring, e.g., polymorphic or allelic variants, and man-made variants. As described below, the preferred algorithms can account for gaps and the like. Preferably, identity exists over a region that is at least about 25 amino acids or nucleotides in length, or more preferably over a region that is 50-100 amino acids or

nucleotides in length.

For sequence comparison, typically one sequence acts as a reference sequence, to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are entered into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. Preferably, default program parameters can be used, or alternative parameters can be designated. The sequence comparison algorithm then calculates the percent sequence identities for the test sequences relative to the reference sequence, based on the program parameters.

A "comparison window", as used herein, includes reference to a segment of contiguous positions selected from the group consisting typically of from 20 to 600, usually about 50 to about 200, more usually about 100 to about 150 in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned. Methods of alignment of sequences for comparison are well-known in the art. Optimal alignment of sequences for comparison can be conducted, e.g., by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482, by the homology alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity method of Pearson and Lipman (1988) Proc. Nat'l. Acad. Sci. USA 85:2444, by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer

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Group, 575 Science Dr., Madison, WI), or by manual alignment and visual inspection (see, e.g., Ausubel, et al. (eds. 1995 and supplements) <u>Current Protocols in Molecular Biology</u>.

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Preferred examples of algorithms that are suitable for determining percent sequence identity and sequence similarity include the BLAST and BLAST 2.0 algorithms, which are described in Altschul, et al. (1977) Nuc. Acids Res. 25:3389-3402 and Altschul, et al. (1990) J. Mol. Biol. 215:403-410. BLAST and BLAST 2.0 are used, with the parameters described herein, to determine percent sequence identity for the nucleic acids and proteins of the invention. Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (http://www.ncbi.nlm.nih.gov/). This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul, et al., supra). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, e.g., for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, an expectation (E) of 10, M=5, N=-4 and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength of 3, and expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915) alignments (B) of 50, expectation (E) of 10, M=5, N=-4, and a comparison of both strands.

The BLAST algorithm also performs a statistical analysis of the similarity between two sequences (see, e.g., Karlin and Altschul (1993) <u>Proc. Nat'l. Acad. Sci. USA</u> 90:5873-5787). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between

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two nucleotide or amino acid sequences would occur by chance. For example, a nucleic acid is considered similar to a reference sequence if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.2, more preferably less than about 0.01, and most preferably less than about 0.001. Log values may be negative large numbers, e.g., 5, 10, 20, 30, 40, 40, 70, 90, 110, 150, 170, etc.

An indication that two nucleic acid sequences are substantially identical is that the polypeptide encoded by the first nucleic acid is immunologically cross reactive with the antibodies raised against the polypeptide encoded by the second nucleic acid. Thus, a polypeptide is typically substantially identical to a second polypeptide, e.g., where the two peptides differ only by conservative substitutions. Another indication that two nucleic acid sequences are substantially identical is that the two molecules or their complements hybridize to each other under stringent conditions. Yet another indication that two nucleic acid sequences are substantially identical is that the same primers can be used to amplify the sequences.

A "host cell" is a naturally occurring cell or a transformed cell that contains an expression vector and supports the replication or expression of the expression vector. Host cells may be cultured cells, explants, cells *in vivo*, and the like. Host cells may be prokaryotic cells such as *E. coli*, or eukaryotic cells such as yeast, insect, amphibian, or mammalian cells such as CHO, HeLa, and the like (see, e.g., the American Type Culture Collection catalog or web site, www.atcc.org).

The terms "isolated," "purified," or "biologically pure" refer to material that is substantially or essentially free from components that normally accompany it as found in its native state. Purity and homogeneity are typically determined using analytical chemistry techniques such as polyacrylamide gel electrophoresis or high performance liquid chromatography. A protein or nucleic acid that is the predominant species present in a preparation is substantially purified. In particular, an isolated nucleic acid is separated from some open reading frames that naturally flank the gene and encode proteins other than protein encoded by the gene. The term "purified" in some embodiments denotes that a nucleic acid or protein gives rise to essentially one band in an electrophoretic gel. Preferably, it means that the nucleic acid or protein is at least about 85% pure, more preferably at least 95% pure, and most preferably at least 99% pure. "Purify" or "purification" in other embodiments means removing at least one contaminant or component from the composition to be purified.

WO 02/086443 PCT/US02/12476 In this sense, purification does not require that the purified compound be homogeneous, e.g., 100% pure.

The terms "polypeptide," "peptide" and "protein" are used interchangeably herein to refer to a polymer of amino acid residues. The terms apply to amino acid polymers in which one or more amino acid residue is an artificial chemical mimetic of a corresponding naturally occurring amino acid, as well as to naturally occurring amino acid polymers, those containing modified residues, and non-naturally occurring amino acid polymer.

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The term "amino acid" refers to naturally occurring and synthetic amino acids, as well as amino acid analogs and amino acid mimetics that function similarly to the naturally occurring amino acids. Naturally occurring amino acids are those encoded by the genetic code, as well as those amino acids that are later modified, e.g., hydroxyproline, γ-carboxyglutamate, and O-phosphoserine. Amino acid analogs refer to compounds that have the same basic chemical structure as a naturally occurring amino acid, e.g., an α carbon that is bound to a hydrogen, a carboxyl group, an amino group, and an R group, e.g., homoserine, norleucine, methionine sulfoxide, methionine methyl sulfonium. Such analogs may have modified R groups (e.g., norleucine) or modified peptide backbones, but retain some basic chemical structure as a naturally occurring amino acid. Amino acid mimetics refer to chemical compounds that have a structure that is different from the general chemical structure of an amino acid, but that function similarly to another amino acid.

Amino acids may be referred to herein by either their commonly known three letter symbols or by the one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Nucleotides, likewise, may be referred to by their commonly accepted single-letter codes.

"Conservatively modified variants" applies to both amino acid and nucleic acid sequences. With respect to particular nucleic acid sequences, conservatively modified variants refers to those nucleic acids which encode identical or essentially identical amino acid sequences, or where the nucleic acid does not encode an amino acid sequence, to essentially identical or associated, e.g., naturally contiguous, sequences. Because of the degeneracy of the genetic code, a large number of functionally identical nucleic acids encode most proteins. For instance, the codons GCA, GCC, GCG, and GCU each encode the amino acid alanine. Thus, at each position where an alanine is specified by a codon, the codon can be altered to another of the corresponding codons described without altering the encoded polypeptide. Such nucleic acid variations are "silent variations," which are one species of

conservatively modified variations. Every nucleic acid sequence herein which encodes a polypeptide also describes silent variations of the nucleic acid. In certain contexts each codon in a nucleic acid (except AUG, which is ordinarily the only codon for methionine, and TGG, which is ordinarily the only codon for tryptophan) can be modified to yield a functionally similar molecule. Accordingly, a silent variation of a nucleic acid which encodes a polypeptide is implicit in a described sequence with respect to the expression product, but not necessarily with respect to actual probe sequences.

As to amino acid sequences, one of skill will recognize that individual substitutions, deletions or additions to a nucleic acid, peptide, polypeptide, or protein sequence which alters, adds or deletes a single amino acid or a small percentage of amino acids in the encoded sequence is a "conservatively modified variant" where the alteration results in the substitution of an amino acid with a chemically similar amino acid. Conservative substitution tables providing functionally similar amino acids are well known in the art. Such conservatively modified variants are in addition to and do not exclude polymorphic variants, interspecies homologs, and alleles of the invention. Typically conservative substitutions include for one another: 1) Alanine (A), Glycine (G); 2) Aspartic acid (D), Glutamic acid (E); 3) Asparagine (N), Glutamine (Q); 4) Arginine (R), Lysine (K); 5) Isoleucine (I), Leucine (L), Methionine (M), Valine (V); 6) Phenylalanine (F), Tyrosine (Y), Tryptophan (W); 7) Serine (S), Threonine (T); and 8) Cysteine (C), Methionine (M) (see, e.g., Creighton, Proteins (1984)).

Macromolecular structures such as polypeptide structures can be described in terms of various levels of organization. For a general discussion of this organization, see, e.g., Alberts, et al. (1994) Molecular Biology of the Cell (3^{rd} ed.) and Cantor and Schimmel (1980) Biophysical Chemistry Part I: The Conformation of Biological Macromolecules. "Primary structure" refers to the amino acid sequence of a particular peptide. "Secondary structure" refers to locally ordered, three dimensional structures within a polypeptide. These structures are commonly known as domains. Domains are portions of a polypeptide that often form a compact unit of the polypeptide and are typically 25 to approximately 500 amino acids long. Typical domains are made up of sections of lesser organization such as stretches of β -sheet and α -helices. "Tertiary structure" refers to the complete three dimensional structure of a polypeptide monomer. "Quaternary structure" refers to the three dimensional structure formed, usually by the noncovalent association of independent tertiary units. Anisotropic terms are also known as energy terms.

"Nucleic acid" or "oligonucleotide" or "polynucleotide" or grammatical equivalents used herein means at least two nucleotides covalently linked together. Oligonucleotides are typically from about 5, 6, 7, 8, 9, 10, 12, 15, 25, 30, 40, 50 or more nucleotides in length, up to about 100 nucleotides in length. Nucleic acids and polynucleotides are a polymers of any length, including longer lengths, e.g., 200, 300, 500, 1000, 2000, 3000, 5000, 7000, 10,000, 5 etc. A nucleic acid of the present invention will generally contain phosphodiester bonds, although in some cases, nucleic acid analogs are included that may have at least one different linkage, e.g., phosphoramidate, phosphorothioate, phosphorodithioate, or Omethylphophoroamidite linkages (see Eckstein (1992) Oligonucleotides and Analogues: A Practical Approach Oxford University Press); and peptide nucleic acid backbones and 10 linkages. Other analog nucleic acids include those with positive backbones; non-ionic backbones, and non-ribose backbones, including those described in U.S. Patent Nos. 5,235,033 and 5,034,506, and Chapters 6 and 7, in Sanghui and Cook, eds. Carbohydrate Modifications in Antisense Research, ASC Symposium Series 580. Nucleic acids containing one or more carbocyclic sugars are also included within one definition of nucleic acids. 15 Modifications of the ribose-phosphate backbone may be done for a variety of reasons, e.g., to increase the stability and half-life of such molecules in physiological environments or as probes on a biochip. Mixtures of naturally occurring nucleic acids and analogs can be made; alternatively, mixtures of different nucleic acid analogs, and mixtures of naturally occurring nucleic acids and analogs may be made. 20

Particularly preferred are peptide nucleic acids (PNA) which includes peptide nucleic acid analogs. These backbones are substantially non-ionic under neutral conditions, in contrast to the highly charged phosphodiester backbone of naturally occurring nucleic acids. This results in two advantages. First, the PNA backbone exhibits improved hybridization kinetics. PNAs have larger changes in the melting temperature (T_m) for mismatched versus perfectly matched basepairs. DNA and RNA typically exhibit a 2-4° C drop in T_m for an internal mismatch. With the non-ionic PNA backbone, the drop is closer to 7-9° C. Similarly, due to their non-ionic nature, hybridization of the bases attached to these backbones is relatively insensitive to salt concentration. In addition, PNAs are not degraded by cellular enzymes, and thus can be more stable.

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The nucleic acids may be single stranded or double stranded, as specified, or contain portions of both double stranded or single stranded sequence. As will be appreciated by those in the art, the depiction of a single strand also defines the sequence of the complementary

strand; thus the sequences described herein also provide the complement of the sequence. The nucleic acid may be DNA, both genomic and cDNA, RNA, or a hybrid, where the nucleic acid may contain combinations of deoxyribo- and ribo-nucleotides, and combinations of bases, including uracil, adenine, thymine, cytosine, guanine, inosine, xanthine hypoxanthine, isocytosine, isoguanine, etc. "Transcript" typically refers to a naturally occurring RNA, e.g., a pre-mRNA, hnRNA, or mRNA. As used herein, the term "nucleoside" includes nucleotides and nucleoside and nucleotide analogs, and modified nucleosides such as amino modified nucleosides. In addition, "nucleoside" includes non-naturally occurring analog structures. Thus, e.g., the individual units of a peptide nucleic acid, each containing a base, are referred to herein as a nucleoside.

A "label" or a "detectable moiety" is a composition detectable by spectroscopic, photochemical, biochemical, immunochemical, physiological, chemical, or other physical means. For example, useful labels include ³²P, fluorescent dyes, electron-dense reagents, enzymes (e.g., as commonly used in an ELISA), biotin, digoxigenin, or haptens and proteins or other entities which can be made detectable, e.g., by incorporating a radiolabel into the peptide or used to detect antibodies specifically reactive with the peptide. The labels may be incorporated into the cancer nucleic acids, proteins, and antibodies. Many methods known in the art for conjugating the antibody to the label may be employed, including those methods described by Hunter, et al. (1962) Nature 144:945; David, et al. (1974) Biochemistry 13:1014-1021; Pain, et al. (1981) J. Immunol. Meth., 40:219-230; and Nygren (1982) J. Histochem. and Cytochem. 30:407-412.

An "effector" or "effector moiety" or "effector component" is a molecule that is bound (or linked, or conjugated), either covalently, through a linker or a chemical bond, or noncovalently, through ionic, van der Waals, electrostatic, or hydrogen bonds, to an antibody. The "effector" can be a variety of molecules including, e.g., detection moieties including radioactive compounds, fluorescent compounds, an enzyme or substrate, tags such as epitope tags, a toxin; activatable moieties, a chemotherapeutic agent; a lipase; an antibiotic; or a radioisotope emitting "hard" e.g., beta radiation.

A "labeled nucleic acid probe or oligonucleotide" is one that is bound, either covalently, through a linker or a chemical bond, or noncovalently, through ionic, van der Waals, electrostatic, or hydrogen bonds to a label such that the presence of the probe may be detected by detecting the presence of the label bound to the probe. Alternatively, method

using high affinity interactions may achieve the same results where one of a pair of binding partners binds to the other, e.g., biotin, streptavidin.

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As used herein a "nucleic acid probe or oligonucleotide" is a nucleic acid capable of binding to a target nucleic acid of complementary sequence through one or more types of chemical bonds, usually through complementary base pairing, e.g., through hydrogen bond formation. As used herein, a probe may include natural (i.e., A, G, C, or T) or modified bases (7-deazaguanosine, inosine, etc.). In addition, the bases in a probe may be joined by a linkage other than a phosphodiester bond, preferably one that does not functionally interfere with hybridization. Thus, e.g., probes may be peptide nucleic acids in which the constituent bases are joined by peptide bonds rather than phosphodiester linkages. Probes may bind target sequences lacking complete complementarity with the probe sequence depending upon the stringency of the hybridization conditions. The probes are preferably directly labeled, e.g., with isotopes, chromophores, lumiphores, chromogens, or indirectly labeled, e.g., with biotin to which a streptavidin complex may later bind. By assaying for the presence or absence of the probe, one can detect the presence or absence of the select sequence or subsequence. Diagnosis or prognosis may be based at the genomic level, or at the level of RNA or protein expression.

The term "recombinant" when used with reference, e.g., to a cell, or nucleic acid, protein, or vector, indicates that the cell, nucleic acid, protein or vector, has been modified by the introduction of a heterologous nucleic acid or protein or the alteration of a native nucleic acid or protein, or that the cell is derived from a cell so modified. Thus, e.g., recombinant cells express genes that are not found within the native (non-recombinant) form of the cell or express native genes that are otherwise abnormally expressed, under expressed or not expressed at all. By the term "recombinant nucleic acid" herein is meant nucleic acid, originally formed in vitro, in general, by the manipulation of nucleic acid, e.g., using polymerases and endonucleases, in a form not normally found in nature. In this manner, operably linkage of different sequences is achieved. Thus an isolated nucleic acid, in a linear form, or an expression vector formed in vitro by ligating DNA molecules that are not normally joined, are both considered recombinant for the purposes of this invention. It is understood that once a recombinant nucleic acid is made and reintroduced into a host cell or organism, it will replicate non-recombinantly, i.e., using the in vivo cellular machinery of the host cell rather than in vitro manipulations; however, such nucleic acids, once produced recombinantly, although subsequently replicated non-recombinantly, are still considered

recombinant for the purposes of the invention. Similarly, a "recombinant protein" is a protein made using recombinant techniques, i.e., through the expression of a recombinant nucleic acid as depicted above.

The term "heterologous" when used with reference to portions of a nucleic acid indicates that the nucleic acid comprises two or more subsequences that are not normally found in the same relationship to each other in nature. For instance, the nucleic acid is typically recombinantly produced, having two or more sequences, e.g., from unrelated genes arranged to make a new functional nucleic acid, e.g., a promoter from one source and a coding region from another source. Similarly, a heterologous protein will often refer to two or more subsequences that are not found in the same relationship to each other in nature (e.g., a fusion protein).

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A "promoter" is typically an array of nucleic acid control sequences that direct transcription of a nucleic acid. As used herein, a promoter includes necessary nucleic acid sequences near the start site of transcription, such as, in the case of a polymerase II type promoter, a TATA element. A promoter also optionally includes distal enhancer or repressor elements, which can be located as much as several thousand base pairs from the start site of transcription. A "constitutive" promoter is a promoter that is active under most environmental and developmental conditions. An "inducible" promoter is a promoter that is active under environmental or developmental regulation. The term "operably linked" refers to a functional linkage between a nucleic acid expression control sequence (such as a promoter, or array of transcription factor binding sites) and a second nucleic acid sequence, e.g., wherein the expression control sequence directs transcription of the nucleic acid corresponding to the second sequence.

An "expression vector" is a nucleic acid construct, generated recombinantly or synthetically, with a series of specified nucleic acid elements that permit transcription of a particular nucleic acid in a host cell. The expression vector can be part of a plasmid, virus, or nucleic acid fragment. Typically, the expression vector includes a nucleic acid to be transcribed in operable linkage to a promoter.

The phrase "selectively (or specifically) hybridizes to" refers to the binding, duplexing, or hybridizing of a molecule selectively to a particular nucleotide sequence under stringent hybridization conditions when that sequence is present in a complex mixture (e.g., total cellular or library DNA or RNA).

The phrase "stringent hybridization conditions" refers to conditions under which a probe will hybridize to its target subsequence, typically in a complex mixture of nucleic acids, but to essentially no other sequences. Stringent conditions are sequence-dependent and will be different in different circumstances. Longer sequences hybridize specifically at higher temperatures. An extensive guide to the hybridization of nucleic acids is found in . 5 "Overview of principles of hybridization and the strategy of nucleic acid assays" in Tijssen (1993) Techniques in Biochemistry and Molecular Biology--Hybridization with Nucleic Probes (vol. 24) Elsevier. Generally, stringent conditions are selected to be about 5-10° C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength pH. The T_m is the temperature (under defined ionic strength, pH, and nucleic concentration) 10 at which 50% of the probes complementary to the target hybridize to the target sequence at equilibrium (as the target sequences are present in excess, at T_m, 50% of the probes are occupied at equilibrium). Stringent conditions will be those in which the salt concentration is less than about 1.0 M sodium ion, typically about 0.01 to 1.0 M sodium ion concentration (or other salts) at pH 7.0 to 8.3 and the temperature is at least about 30° C for short probes (e.g., 15 10 to 50 nucleotides) and at least about 60° C for long probes (e.g., greater than 50 nucleotides). Stringent conditions may also be achieved with the addition of destabilizing agents such as formamide. For selective or specific hybridization, a positive signal is typically at least two times background, preferably 10 times background hybridization. Exemplary stringent hybridization conditions are often: 50% formamide, 5x SSC, and 1% 20 SDS, incubating at 42° C, or, 5x SSC, 1% SDS, incubating at 65° C, with wash in 0.2x SSC, and 0.1% SDS at 65° C. For PCR, a temperature of about 36° C is typical for low stringency amplification, although annealing temperatures may vary between about 32° C and 48° C depending on primer length. For high stringency PCR amplification, a temperature of about 25 62° C is typical, although high stringency annealing temperatures can range from about 50° C to about 65° C, depending on the primer length and specificity. Typical cycle conditions for both high and low stringency amplifications include a denaturation phase of 90° C - 95° C for 0.5 - 2 min., an annealing phase lasting 0.5 - 2 min., and an extension phase of about 72° C for 1 - 2 min. Protocols and guidelines for low and high stringency amplification reactions are provided, e.g., in Innis, et al.(1990) PCR Protocols, A Guide to Methods and 30 Applications.

Nucleic acids that do not hybridize to each other under stringent conditions are still substantially identical if the polypeptides which they encode are substantially identical. This

occurs, e.g., when a copy of a nucleic acid is created using the maximum codon degeneracy permitted by the genetic code. In such cases, the nucleic acids typically hybridize under moderately stringent hybridization conditions. Exemplary "moderately stringent hybridization conditions" include a hybridization in a buffer of 40% formamide, 1 M NaCl, 1% SDS at 37° C, and a wash in 1X SSC at 45° C. A positive hybridization is at least twice background. Alternative hybridization and wash conditions can be utilized to provide conditions of similar stringency. Additional guidelines for determining hybridization parameters are provided in numerous reference, e.g., Ausubel, et al. (ed.) <u>Current Protocols in</u>

Molecular Biology Lippincott.

The phrase "functional effects" in the context of assays for testing compounds that modulate activity of a lung cancer protein includes the determination of a parameter that is indirectly or directly under the influence of the lung cancer protein or nucleic acid, e.g., a physiological, enzymatic, functional, physical, or chemical effect, such as the ability to decrease lung cancer. It includes ligand binding activity; cell viability, cell growth on soft agar; anchorage dependence; contact inhibition and density limitation of growth; cellular proliferation; cellular transformation; growth factor or serum dependence; tumor specific marker levels; invasiveness into Matrigel; tumor growth and metastasis *in vivo*; mRNA and protein expression in cells undergoing metastasis, and other characteristics of lung cancer cells. "Functional effects" include *in vitro*, *in vivo*, and *ex vivo* activities.

By "determining the functional effect" is meant assaying for a compound that increases or decreases a parameter that is indirectly or directly under the influence of a lung cancer protein sequence, e.g., physiological, functional, enzymatic, physical, or chemical effects. Such functional effects can be measured by many means known to those skilled in the art, e.g., changes in spectroscopic characteristics (e.g., fluorescence, absorbance, refractive index), hydrodynamic (e.g., shape), chromatographic, or solubility properties for the protein, measuring inducible markers or transcriptional activation of the lung cancer protein; measuring binding activity or binding assays, e.g., binding to antibodies or other ligands, and measuring cellular proliferation. Determination of the functional effect of a compound on lung cancer can also be performed using lung cancer assays known to those of skill in the art such as an *in vitro* assays, e.g., cell growth on soft agar; anchorage dependence; contact inhibition and density limitation of growth; cellular proliferation; cellular transformation; growth factor or serum dependence; tumor specific marker levels; invasiveness into Matrigel; tumor growth and metastasis *in vivo*; mRNA and protein

expression in cells undergoing metastasis, and other characteristics of lung cancer cells. The functional effects can be evaluated by many means known to those skilled in the art, e.g., microscopy for quantitative or qualitative measures of alterations in morphological features, measurement of changes in RNA or protein levels for lung cancer-associated sequences, measurement of RNA stability, identification of downstream or reporter gene expression (CAT, luciferase, β -gal, GFP, and the like), e.g., via chemiluminescence, fluorescence, colorimetric reactions, antibody binding, inducible markers, and ligand binding assays.

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"Inhibitors", "activators", and "modulators" of lung cancer polynucleotide and polypeptide sequences are used to refer to activating, inhibitory, or modulating molecules or compounds identified using in vitro and in vivo assays of lung cancer polynucleotide and polypeptide sequences. Inhibitors are compounds that, e.g., bind to, partially or totally block activity, decrease, prevent, delay activation, inactivate, desensitize, or down regulate the activity or expression of lung cancer proteins, e.g., antagonists. Antisense or inhibitory nucleic acids may seem to inhibit expression and subsequent function of the protein. "Activators" are compounds that increase, open, activate, facilitate, enhance activation, sensitize, agonize, or up regulate lung cancer protein activity. Inhibitors, activators, or modulators also include genetically modified versions of lung cancer proteins, e.g., versions with altered activity, as well as naturally occurring and synthetic ligands, antagonists, agonists, antibodies, small chemical molecules and the like. Such assays for inhibitors and activators include, e.g., expressing the lung cancer protein in vitro, in cells, or cell membranes, applying putative modulator compounds, and then determining the functional effects on activity, as described above. Activators and inhibitors of lung cancer can also be identified by incubating lung cancer cells with the test compound and determining increases or decreases in the expression of 1 or more lung cancer proteins, e.g., 1, 2, 3, 4, 5, 10, 15, 20, 25, 30, 40, 50 or more lung cancer proteins, such as lung cancer proteins encoded by the sequences set out in Tables 1A-16.

Samples or assays comprising lung cancer proteins that are treated with a potential activator, inhibitor, or modulator are compared to control samples without the inhibitor, activator, or modulator to examine the extent of inhibition. Control samples (untreated with inhibitors) are assigned a relative protein activity value of 100%. Inhibition of a polypeptide is achieved when the activity value relative to the control is about 80%, preferably 50%, more preferably 25-0%. Activation of a lung cancer polypeptide is achieved when the activity value relative to the control (untreated with activators) is 110%, more preferably 150%, more

WO 02/086443 PCT/US02/12476 preferably 200-500% (i.e., two to five fold higher relative to the control), more preferably 1000-3000% higher.

The phrase "changes in cell growth" refers to any change in cell growth and proliferation characteristics *in vitro* or *in vivo*, such as cell viability, formation of foci, anchorage independence, semi-solid or soft agar growth, changes in contact inhibition and density limitation of growth, loss of growth factor or serum requirements, changes in cell morphology, gaining or losing immortalization, gaining or losing tumor specific markers, ability to form or suppress tumors when injected into suitable animal hosts, and/or immortalization of the cell. See, e.g., Freshney (1994) <u>Culture of Animal Cells a Manual of Basic Technique</u> pp. 231-241 (3rd ed.).

"Tumor cell" refers to precancerous, cancerous, and normal cells in a tumor.

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"Cancer cells," "transformed" cells, or "transformation" in tissue culture, refers to spontaneous or induced phenotypic changes that do not necessarily involve the uptake of new genetic material. Although transformation can arise from infection with a transforming virus and incorporation of new genomic DNA, or uptake of exogenous DNA, it can also arise spontaneously or following exposure to a carcinogen, thereby mutating an endogenous gene. Transformation is associated with phenotypic changes, such as immortalization of cells, aberrant growth control, nonmorphological changes, and/or malignancy (see, Freshney (1994) Culture of Animal Cells a Manual of Basic Technique (3rd ed.)).

"Antibody" refers to a polypeptide comprising a framework region from an immunoglobulin gene or fragments thereof that specifically binds and recognizes an antigen. The recognized immunoglobulin genes include the kappa, lambda, alpha, gamma, delta, epsilon, and mu constant region genes, as well as the myriad immunoglobulin variable region genes. Light chains are classified as either kappa or lambda. Heavy chains are classified as gamma, mu, alpha, delta, or epsilon, which in turn define the immunoglobulin classes, IgG, IgM, IgA, IgD, and IgE, respectively. Typically, the antigen-binding region of an antibody or its functional equivalent will be most critical in specificity and affinity of binding. See Paul, Fundamental Immunology.

An exemplary immunoglobulin (antibody) structural unit comprises a tetramer. Each tetramer is composed of two identical pairs of polypeptide chains, each pair having one "light" (about 25 kD) and one "heavy" chain (about 50-70 kD). The N-terminus of each chain defines a variable region of about 100 to 110 or more amino acids primarily responsible

WO 02/086443 PCT/US02/12476 for antigen recognition. The terms variable light chain (V_L) and variable heavy chain (V_H) refer to these light and heavy chains respectively.

Antibodies exist, e.g., as intact immunoglobulins or as a number of well-characterized fragments produced by digestion with various peptidases. Thus, e.g., pepsin digests an antibody below the disulfide linkages in the hinge region to produce F(ab)'₂, a dimer of Fab which itself is a light chain joined to V_H-C_H1 by a disulfide bond. The F(ab)'₂ may be reduced under mild conditions to break the disulfide linkage in the hinge region, thereby converting the F(ab)'₂ dimer into an Fab' monomer. The Fab' monomer is essentially Fab with part of the hinge region (see Paul (ed. 1999) Fundamental Immunology (4th ed.). While various antibody fragments are defined in terms of the digestion of an intact antibody, one of skill will appreciate that such fragments may be synthesized *de novo* either chemically or by using recombinant DNA methodology. Thus, the term antibody, as used herein, also includes antibody fragments either produced by the modification of whole antibodies, or those synthesized *de novo* using recombinant DNA methodologies (e.g., single chain Fv) or those identified using phage display libraries (see, e.g., McCafferty, et al. (1990) Nature 348:552-554).

For preparation of antibodies, e.g., recombinant, monoclonal, or polyclonal antibodies, many technique known in the art can be used (see, e.g., Kohler and Milstein (1975) Nature 256:495-497; Kozbor, et al. (1983) Immunology Today 4:72; Cole, et al. (1985), pp. 77-96 in Monoclonal Antibodies and Cancer Therapy; Coligan (1991 and supplements) Current Protocols in Immunology; Harlow and Lane (1988) Antibodies, A Laboratory Manual; and Goding (1986) Monoclonal Antibodies: Principles and Practice (2d ed.)). Techniques for the production of single chain antibodies (U.S. Patent 4,946,778) can be adapted to produce antibodies to polypeptides of this invention. Also, transgenic mice, or other organisms such as other mammals, may be used to express humanized antibodies. Alternatively, phage display technology can be used to identify antibodies and heteromeric Fab fragments that specifically bind to selected antigens (see, e.g., McCafferty, et al. (1990) Nature 348:552-554; Marks, et al. (1992) Biotechnology 10:779-783).

A "chimeric antibody" is an antibody molecule in which, e.g, (a) the constant region, or a portion thereof, is altered, replaced, or exchanged so that the antigen binding site (variable region) is linked to a constant region of a different or altered class, effector function, and/or species, or an entirely different molecule which confers new properties to the chimeric antibody, e.g., an enzyme, toxin, hormone, growth factor, drug, etc.; or (b) the

variable region, or a portion thereof, is altered, replaced, or exchanged with a variable region having a different or altered antigen specificity.

Identification of lung cancer-associated sequences

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In one aspect, the expression levels of genes are determined in different patient samples for which diagnosis information is desired, to provide expression profiles. An expression profile of a particular sample is essentially a "fingerprint" of the state of the sample; while two states may have any particular gene similarly expressed, the evaluation of a number of genes simultaneously allows the generation of a gene expression profile that is characteristic of the state of the cell. That is, normal tissue may be distinguished from cancerous or metastatic cancerous tissue, or metastatic cancerous tissue can be compared with tissue from surviving cancer patients. By comparing expression profiles of tissue in known different lung cancer states, information regarding which genes are important (including both up- and down-regulation of genes) in each of these states is obtained. Molecular profiling may distinguish subtypes of a currently collective disease designation, e.g., different forms of lung cancer (chronic disease, adenocarcinoma, etc.)

The identification of sequences that are differentially expressed in lung cancer versus non-lung cancer tissue allows the use of this information in a number of ways. For example, a particular treatment regime may be evaluated: does a chemotherapeutic drug act to downregulate lung cancer, and thus tumor growth or recurrence, in a particular patient. Alternatively, a treatment step may induce other markers which may be used as targets to destroy tumor cells. Similarly, diagnosis and treatment outcomes may be done or confirmed by comparing patient samples with the known expression profiles. Malignant diseasemay be compared to non-malignant conditions. Metastatic tissue can also be analyzed to determine the stage of lung cancer in the tissue, or origin of primary tumor, e.g., metastasis from a remote primary site. Furthermore, these gene expression profiles (or individual genes) allow screening of drug candidates with an eye to mimicking or altering a particular expression profile; e.g., screening can be done for drugs that suppress the lung cancer expression profile. This may be done by making biochips comprising sets of the important lung cancer genes, which can then be used in these screens. PCR methods may be applied with selected primer pairs, and analysis may be of RNA or of genomic sequences. These methods can also be done on the protein basis; that is, protein expression levels of the lung cancer proteins can be evaluated for diagnostic purposes or to screen candidate agents. In addition, the lung cancer

nucleic acid sequences can be administered for gene therapy purposes, including the administration of antisense nucleic acids, or the lung cancer proteins (including antibodies and other modulators thereof) administered as therapeutic drugs or as protein or DNA vaccines.

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Thus the present invention provides nucleic acid and protein sequences that are differentially expressed in lung cancer relative to normal tissues and/or non-malignant lung disease, or in different types of lung disease, herein termed "lung cancer sequences." As outlined below, lung cancer sequences include those that are up-regulated (i.e., expressed at a higher level) in lung cancer, as well as those that are down-regulated (i.e., expressed at a lower level). In a preferred embodiment, the lung cancer sequences are from humans; however, as will be appreciated by those in the art, lung cancer sequences from other organisms may be useful in animal models of disease and drug evaluation; thus, other lung cancer sequences are provided, from vertebrates, including mammals, including rodents (rats, mice, hamsters, guinea pigs, etc.), primates, farm animals (including sheep, goats, pigs, cows, horses, etc.) and pets (dogs, cats, etc.). Lung cancer sequences from other organisms may be obtained using the techniques outlined below.

Lung cancer sequences can include both nucleic acid and amino acid sequences. As will be appreciated by those in the art and is more fully outlined below, lung cancer nucleic acid sequences are useful in a variety of applications, including diagnostic applications, which will detect naturally occurring nucleic acids, as well as screening applications; e.g., biochips comprising nucleic acid probes or PCR microtiter plates with selected probes to the lung cancer sequences can be generated.

A lung cancer sequence can be initially identified by substantial nucleic acid and/or amino acid sequence homology to the lung cancer sequences outlined herein. Such homology can be based upon the overall nucleic acid or amino acid sequence, and is generally determined as outlined below, e.g., using homology programs or hybridization conditions.

For identifying lung cancer-associated sequences, the lung cancer screen typically includes comparing genes identified in different tissues, e.g., normal and cancerous tissues, cancer and non-malignant conditions, non-malignant conditions and normal tissues, or tumor tissue samples from patients who have metastatic disease vs. non metastatic tissue. Other suitable tissue comparisons include comparing lung cancer samples with metastatic cancer samples from other cancers, such as, breast, other gastrointestinal cancers, prostate, ovarian,

etc. Samples of, non metastatic disease tissue and tissue undergoing metastasis are applied to biochips comprising nucleic acid probes. The samples are first microdissected, if applicable, and treated as is known in the art for the preparation of mRNA. Suitable biochips are commercially available, e.g., from Affymetrix, Santa Clara, CA. Gene expression profiles as described herein are generated and the data analyzed.

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In one embodiment, the genes showing changes in expression as between normal and disease states are compared to genes expressed in other normal tissues, preferably normal lung, but also including, and not limited to colon, heart, brain, liver, breast, kidney, muscle, prostate, small intestine, large intestine, spleen, bone, and/or placenta. In a preferred embodiment, those genes identified during the lung cancer screen that are expressed in significant amounts in other tissues (e.g., essential organs) are removed from the profile, although in some embodiments, this is not necessary (e.g., where organs may be dispensible at a later stage of life). That is, when screening for drugs, it is usually preferable that the target expression be disease specific, to minimize possible side effects on other organs.

In a preferred embodiment, lung cancer sequences are those that are up-regulated in lung cancer; that is, the expression of these genes is higher in cancerous tissue than in normal lung or other tissue. "Up-regulation" as used herein means, when the ratio is presented as a number greater than one, that the ratio is greater than one, preferably 1.5 or greater, more preferably 2.0 or greater. Another embodiment is directed to sequences up-regulated in nonmalignant conditions relative to normal. Unigene cluster identification numbers and accession numbers herein are for the GenBank sequence database and the sequences of the accession numbers are hereby expressly incorporated by reference. GenBank is known in the art, see, e.g., Benson, DA, et al (1998) Nucleic Acids Research 26:1-7 and http://www.ncbi.nlm.nih.gov/. Sequences are also available in other databases, e.g., European Molecular Biology Laboratory (EMBL) and DNA Database of Japan (DDBJ). Another embodiment is directed to sequences up-regulated in non-malignant conditions relative to normal. In some situations, the sequences may be derived from assembly of available sequences or be predicted from genomic DNA using exon prediction algorithms, such as FGENESH (Salamov and Solovyev (2000) Genome Res. 10:516-522). In other situations, sequences have been derived from cloning and sequencing of isolated nucleic acids.

In another preferred embodiment, lung cancer sequences are those that are downregulated in the lung cancer; that is, the expression of these genes is lower in cancerous tissue

or normal lung or other tissue. "Down-regulation" as used herein means, when the ratio is presented as a number greater than one, that the ratio is greater than one, preferably 1.5 or greater, more preferably 2.0 or greater, or, when the ratio is presented as a number less than one, that the ratio is less than one, preferably 0.5 or less, more preferably 0.25 or less.

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Informatics

The ability to identify genes that are over or under expressed in lung cancer can additionally provide high-resolution, high-sensitivity datasets which can be used in the areas of diagnostics, therapeutics, drug development, pharmacogenetics, protein structure, biosensor development, and other related areas. For example, the expression profiles can be used in diagnostic or prognostic evaluation of patients with lung cancer. Or as another example, subcellular toxicological information can be generated to better direct drug structure and activity correlation (see Anderson (1998) Pharmaceutical Proteomics: Targets,

Mechanism, and Function, paper presented at the IBC Proteomics conference, Coronado, CA (June 11-12, 1998)). Subcellular toxicological information can also be utilized in a biological sensor device to predict the likely toxicological effect of chemical exposures and likely tolerable exposure thresholds (see U.S. Patent No. 5,811,231). Similar advantages accrue from datasets relevant to other biomolecules and bioactive agents (e.g., nucleic acids, saccharides, lipids, drugs, and the like).

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Thus, in another embodiment, the present invention provides a database that includes at least one set of assay data. The data contained in the database is acquired, e.g., using array analysis either singly or in a library format. The database can be in a form in which data can be maintained and transmitted, but is preferably an electronic database. The electronic database of the invention can be maintained on any electronic device allowing for the storage of and access to the database, such as a personal computer, but is preferably distributed on a wide area network, such as the World Wide Web.

The focus of the present section on databases that include peptide sequence data is for clarity of illustration only. It will be apparent to those of skill in the art that similar databases can be assembled for assay data acquired using an assay of the invention.

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The compositions and methods for identifying and/or quantitating the relative and/or absolute abundance of a variety of molecular and macromolecular species from a biological sample representing lung cancer, i.e., the identification of lung cancer-associated sequences described herein, provide an abundance of information, which can be correlated with

pathological conditions, predisposition to disease, drug testing, therapeutic monitoring, genedisease causal linkages, identification of correlates of immunity and physiological status, among others. Although the data generated from the assays of the invention is suited for manual review and analysis, in a preferred embodiment, data processing using high-speed computers is utilized.

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An array of methods for indexing and retrieving biomolecular information is known in the art. For example, U.S. Patents 6,023,659 and 5,966,712 disclose a relational database system for storing biomolecular sequence information in a manner that allows sequences to be catalogued and searched according to one or more protein function hierarchies. U.S. Patent 5,953,727 discloses a relational database having sequence records containing information in a format that allows a collection of partial-length DNA sequences to be catalogued and searched according to association with one or more sequencing projects for obtaining full-length sequences from the collection of partial length sequences. U.S. Patent 5,706,498 discloses a gene database retrieval system for making a retrieval of a gene sequence similar to a sequence data item in a gene database based on the degree of similarity between a key sequence and a target sequence. U.S. Patent 5,538,897 discloses a method using mass spectroscopy fragmentation patterns of peptides to identify amino acid sequences in computer databases by comparison of predicted mass spectra with experimentally-derived mass spectra using a closeness-of-fit measure. U.S. Patent 5,926,818 discloses a multidimensional database comprising a functionality for multi-dimensional data analysis described as on-line analytical processing (OLAP), which entails the consolidation of projected and actual data according to more than one consolidation path or dimension. U.S. Patent 5,295,261 reports a hybrid database structure in which the fields of each database record are divided into two classes, navigational and informational data, with navigational fields stored in a hierarchical topological map which can be viewed as a tree structure or as the merger of two or more such tree structures.

See also Mount, et al. (2001) Bioinformatics; Durbin, et al. (eds., 1999) Biological

Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids (; Baxevanis and

Oeullette (eds., 1998) Bioinformatics: A Practical Guide to the Analysis of Genes and

Proteins); Rashidi and Buehler (1999) Bioinformatics: Basic Applications in Biological

Science and Medicine; Setubal, et al. (eds 1997) Introduction to Computational Molecular

Biology; Misener and Krawetz (eds, 2000) Bioinformatics: Methods and Protocols; Higgins and Taylor (eds., 2000) Bioinformatics: Sequence, Structure, and Databanks: A Practical

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Approach; Brown (2001) Bioinformatics: A Biologist's Guide to Biocomputing and the

Internet; Han and Kamber (2000) Data Mining: Concepts and Techniques (2000); and

Waterman (1995) Introduction to Computational Biology: Maps, Sequences, and Genomes.

The present invention provides a computer database comprising a computer and software for storing in computer-retrievable form assay data records cross-tabulated, e.g., with data specifying the source of the target-containing sample from which each sequence specificity record was obtained.

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In an exemplary embodiment, at least one of the sources of target-containing sample is from a control tissue sample known to be free of pathological disorders. In a variation, at least one of the sources is a known pathological tissue specimen, e.g., a neoplastic lesion or another tissue specimen to be analyzed for lung cancer. In another variation, the assay records cross-tabulate one or more of the following parameters for each target species in a sample: (1) a unique identification code, which can include, e.g., a target molecular structure and/or characteristic separation coordinate (e.g., electrophoretic coordinates); (2) sample source; and (3) absolute and/or relative quantity of the target species present in the sample.

The invention also provides for the storage and retrieval of a collection of target data in a computer data storage apparatus, which can include magnetic disks, optical disks, magneto-optical disks, DRAM, SRAM, SGRAM, SDRAM, RDRAM, DDR RAM, magnetic bubble memory devices, and other data storage devices, including CPU registers and on-CPU data storage arrays. Typically, the target data records are stored as a bit pattern in an array of magnetic domains on a magnetizable medium or as an array of charge states or transistor gate states, such as an array of cells in a DRAM device (e.g., each cell comprised of a transistor and a charge storage area, which may be on the transistor). In one embodiment, the invention provides such storage devices, and computer systems built therewith, comprising a bit pattern encoding a protein expression fingerprint record comprising unique identifiers for at least 10 target data records cross-tabulated with target source.

When the target is a peptide or nucleic acid, the invention preferably provides a method for identifying related peptide or nucleic acid sequences, comprising performing a computerized comparison between a peptide or nucleic acid sequence assay record stored in or retrieved from a computer storage device or database and at least one other sequence. The comparison can include a sequence analysis or comparison algorithm or computer program embodiment thereof (e.g., FASTA, TFASTA, GAP, BESTFIT) and/or the comparison may

be of the relative amount of a peptide or nucleic acid sequence in a pool of sequences determined from a polypeptide or nucleic acid sample of a specimen.

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The invention also preferably provides a magnetic disk, such as an IBM-compatible (DOS, Windows, Windows95/98/2000, Windows NT, OS/2) or other format (e.g., Linux, SunOS, Solaris, AIX, SCO Unix, VMS, MV, Macintosh, etc.) floppy diskette or hard (fixed, Winchester) disk drive, comprising a bit pattern encoding data from an assay of the invention in a file format suitable for retrieval and processing in a computerized sequence analysis, comparison, or relative quantitation method.

The invention also provides a network, comprising a plurality of computing devices linked via a data link, such as an Ethernet cable (coax or 10BaseT), telephone line, ISDN line, wireless network, optical fiber, or other suitable signal transmission medium, whereby at least one network device (e.g., computer, disk array, etc.) comprises a pattern of magnetic domains (e.g., magnetic disk) and/or charge domains (e.g., an array of DRAM cells) composing a bit pattern encoding data acquired from an assay of the invention.

The invention also provides a method for transmitting assay data that includes generating an electronic signal on an electronic communications device, such as a modem, ISDN terminal adapter, DSL, cable modem, ATM switch, or the like, wherein the signal includes (in native or encrypted format) a bit pattern encoding data from an assay or a database comprising a plurality of assay results obtained by the method of the invention.

In a preferred embodiment, the invention provides a computer system for comparing a query target to a database containing an array of data structures, such as an assay result obtained by the method of the invention, and ranking database targets based on the degree of identity and gap weight to the target data. A central processor is preferably initialized to load and execute the computer program for alignment and/or comparison of the assay results. Data for a query target is entered into the central processor via an I/O device. Execution of the computer program results in the central processor retrieving the assay data from the data file, which comprises a binary description of an assay result.

The target data or record and the computer program can be transferred to secondary memory, which is typically random access memory (e.g., DRAM, SRAM, SGRAM, or SDRAM). Targets are ranked according to the degree of correspondence between a selected assay characteristic (e.g., binding to a selected affinity moiety) and the same characteristic of the query target and results are output via an I/O device. For example, a central processor can be a conventional computer (e.g., Intel Pentium, PowerPC, Alpha, PA-8000, SPARC,

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MIPS 4400, MIPS 10000, VAX, etc.); a program can be a commercial or public domain molecular biology software package (e.g., UWGCG Sequence Analysis Software, Darwin); a data file can be an optical or magnetic disk, a data server, a memory device (e.g., DRAM, SRAM, SGRAM, SDRAM, EPROM, bubble memory, flash memory, etc.); an I/O device can be a terminal comprising a video display and a keyboard, a modem, an ISDN terminal adapter, an Ethernet port, a punched card reader, a magnetic strip reader, or other suitable I/O device.

The invention also preferably provides the use of a computer system, such as that described above, which comprises: (1) a computer; (2) a stored bit pattern encoding a collection of peptide sequence specificity records obtained by the methods of the invention, which may be stored in the computer; (3) a comparison target, such as a query target; and (4) a program for alignment and comparison, typically with rank-ordering of comparison results on the basis of computed similarity values.

Characteristics of lung cancer-associated proteins

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Lung cancer proteins of the present invention may be classified as secreted proteins, transmembrane proteins or intracellular proteins. In one embodiment, the lung cancer protein is an intracellular protein. Intracellular proteins may be found in the cytoplasm and/or in the nucleus. Intracellular proteins are involved in all aspects of cellular function and replication (including, e.g., signaling pathways); aberrant expression of such proteins often results in unregulated or disregulated cellular processes (see, e.g., Alberts (ed. 1994) Molecular Biology of the Cell (3d ed.). For example, many intracellular proteins have enzymatic activity such as protein kinase activity, protein phosphatase activity, protease activity, nucleotide cyclase activity, polymerase activity and the like. Intracellular proteins also serve as docking proteins that are involved in organizing complexes of proteins, or targeting proteins to various subcellular localizations, and are involved in maintaining the structural integrity of organelles.

An increasingly appreciated concept in characterizing proteins is the presence in the proteins of one or more structural motifs for which defined functions have been attributed. In addition to the highly conserved sequences found in the enzymatic domain of proteins, highly conserved sequences have been identified in proteins that are involved in protein-protein interaction. For example, Src-homology-2 (SH2) domains bind tyrosine-phosphorylated targets in a sequence dependent manner. PTB domains, which are distinct from SH2

WO 02/086443 PCT/US02/12476 domains, also bind tyrosine phosphorylated targets. SH3 domains bind to proline-rich

targets. In addition, PH domains, tetratricopeptide repeats and WD domains to name only a few, have been shown to mediate protein-protein interactions. Some of these may also be involved in binding to phospholipids or other second messengers. As will be appreciated by one of ordinary skill in the art, these motifs can be identified on the basis of amino acid sequence; thus, an analysis of the sequence of proteins may provide insight into both the enzymatic potential of the molecule and/or molecules with which the protein may associate. One useful database is Pfam (protein families), which is a large collection of multiple sequence alignments and hidden Markov models covering many common protein domains. Versions are available via the internet from Washington University in St. Louis, the Sanger Center in England, and the Karolinska Institute in Sweden (see, e.g., Bateman, et al (2000) Nuc. Acids Res. 28:263-266; Sonnhammer, et al. (1997) Proteins 28:405-420; Bateman, et al. (1999) Nuc. Acids Res. 27:260-262; and Sonnhammer, et al. (1998) Nuc. Acids Res. 26:320-322).

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In another embodiment, the lung cancer sequences are transmembrane proteins. Transmembrane proteins are molecules that span a phospholipid bilayer of a cell. They may have an intracellular domain, an extracellular domain, or both. The intracellular domains of such proteins may have a number of functions including those already described for intracellular proteins. For example, the intracellular domain may have enzymatic activity and/or may serve as a binding site for additional proteins. Frequently the intracellular domain of transmembrane proteins serves both roles. For example certain receptor tyrosine kinases have both protein kinase activity and SH2 domains. In addition, autophosphorylation of tyrosines on the receptor molecule itself, creates binding sites for additional SH2 domain containing proteins.

Transmembrane proteins may contain from one to many transmembrane domains. For example, receptor tyrosine kinases, certain cytokine receptors, receptor guanylyl cyclases and receptor serine/threonine protein kinases contain a single transmembrane domain. However, various other proteins including channels, pumps, and adenylyl cyclases contain numerous transmembrane domains. Many important cell surface receptors such as G protein coupled receptors (GPCRs) are classified as "seven transmembrane domain" proteins, as they contain 7 membrane spanning regions. Characteristics of transmembrane domains include approximately 17 consecutive hydrophobic amino acids that may be followed by charged amino acids. Therefore, upon analysis of the amino acid sequence of a particular protein, the

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localization and number of transmembrane domains within the protein may be predicted (see, e.g., PSORT web site http://psort.nibb.ac.jp/).

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The extracellular domains of transmembrane proteins are diverse; however, conserved motifs are found repeatedly among various extracellular domains. Conserved structure and/or functions have been ascribed to different extracellular motifs. Many extracellular domains are involved in binding to other molecules. In one aspect, extracellular domains are found on receptors. Factors that bind the receptor domain include circulating ligands, which may be peptides, proteins, or small molecules such as adenosine and the like. For example, growth factors such as EGF, FGF, and PDGF are circulating growth factors that bind to their cognate receptors to initiate a variety of cellular responses. Other factors include cytokines, mitogenic factors, hormones, neurotrophic factors and the like. Extracellular domains also bind to cell-associated molecules. In this respect, they may mediate cell-cell interactions. Cell-associated ligands can be tethered to the cell, e.g., via a glycosylphosphatidylinositol (GPI) anchor, or may themselves be transmembrane proteins. Extracellular domains may also associate with the extracellular matrix and contribute to the maintenance of the cell structure.

Lung cancer proteins that are transmembrane are particularly preferred in the present invention as they are readily accessible targets for extracellular immunotherapeutics, as are described herein. In addition, as outlined below, transmembrane proteins can be also useful in imaging modalities. Antibodies may be used to label such readily accessible proteins in situ or in histological analysis. Alternatively, antibodies can also label intracellular proteins, in which case analytical samples are typically permeablized to provide access to intracellular proteins. In addition, some membrane proteins can be processed to release a soluble protein, or to expose a residual fragment. Released soluble proteins may be useful diagnostic markers, processed residual protein fragments may be useful lung markers of disease.

It will also be appreciated by those in the art that a transmembrane protein can be made soluble by removing transmembrane sequences, e.g., through recombinant methods. Furthermore, transmembrane proteins that have been made soluble can be made to be secreted through recombinant means by adding an appropriate signal sequence.

In another embodiment, the lung cancer proteins are secreted proteins; the secretion of which can be either constitutive or regulated. These proteins may have a signal peptide or signal sequence that targets the molecule to the secretory pathway. Secreted proteins are involved in numerous physiological events; e.g., if circulating, they often serve to transmit

signals to various other cell types. The secreted protein may function in an autocrine manner (acting on the cell that secreted the factor), a paracrine manner (acting on cells in close proximity to the cell that secreted the factor), an endocrine manner (acting on cells at a distance, e.g., secretion into the blood stream), or exocrine (secretion, e.g., through a duct or to adjacent epithelial surface as sweat glands, sebaceous glands, pancreatic ducts, lacrimal glands, mammary glands, sax producing glands of the ear, etc.). Thus secreted molecules often find use in modulating or altering numerous aspects of physiology. Lung cancer proteins that are secreted proteins are particularly preferred in the present invention as they serve as good targets for diagnostic markers, e.g., for blood, plasma, serum, or stool tests. Those which are enzymes may be antibody or small molecule targets. Others may be useful as vaccine targets, e.g., via CTL mechanisms.

Use of lung cancer nucleic acids

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As described above, lung cancer sequence is initially identified by substantial nucleic acid and/or amino acid sequence homology or linkage to the lung cancer sequences outlined herein. Such homology can be based upon the overall nucleic acid or amino acid sequence, and is generally determined as outlined below, using either homology programs or hybridization conditions. Typically, linked sequences on a mRNA are found on the same molecule.

The lung cancer nucleic acid sequences of the invention, e.g., the sequences in Tables 1A-16, can be fragments of larger genes, i.e., they are nucleic acid segments. "Genes" in this context includes coding regions, non-coding regions, and mixtures of coding and non-coding regions. Accordingly, as will be appreciated by those in the art, using the sequences provided herein, extended sequences, in either direction, of the lung cancer genes can be obtained, using techniques well known in the art for cloning either longer sequences or the full length sequences; see Ausubel, et al., *supra*. Much can be done by informatics and many sequences can be clustered to include multiple sequences corresponding to a single gene, e.g., systems such as UniGene (see, http://www.ncbi.nlm.nih.gov/UniGene/).

Once a lung cancer nucleic acid is identified, it can be cloned and, if necessary, its constituent parts recombined to form the entire lung cancer nucleic acid coding regions or the entire mRNA sequence. Once isolated from its natural source, e.g., contained within a plasmid or other vector or excised therefrom as a linear nucleic acid segment, the recombinant lung cancer nucleic acid can be further-used as a probe to identify and isolate

other lung cancer nucleic acids, e.g., extended coding regions. It can also be used as a "precursor" nucleic acid to make modified or variant lung cancer nucleic acids and proteins.

The lung cancer nucleic acids of the present invention are used in several ways. In a first embodiment, nucleic acid probes to the lung cancer nucleic acids are made and attached to biochips to be used in screening and diagnostic methods, as outlined below, or for administration, e.g., for gene therapy, RNAi, vaccine, and/or antisense applications. Alternatively, the lung cancer nucleic acids that include coding regions of lung cancer proteins can be put into expression vectors for the expression of lung cancer proteins, again for screening purposes or for administration to a patient.

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In a preferred embodiment, nucleic acid probes to lung cancer nucleic acids (both the nucleic acid sequences outlined in the figures and/or the complements thereof) are made. The nucleic acid probes attached to the biochip are designed to be substantially complementary to the lung cancer nucleic acids, i.e., the target sequence (either the target sequence of the sample or to other probe sequences, e.g., in sandwich assays), such that hybridization of the target sequence and the probes of the present invention occurs. As outlined below, this complementarity need not be perfect; there may be any number of base pair mismatches which will interfere with hybridization between the target sequence and the single stranded nucleic acids of the present invention. However, if the number of mutations is so great that no hybridization can occur under even the least stringent of hybridization conditions, the sequence is not a complementary target sequence. Thus, by "substantially complementary" herein is meant that the probes are sufficiently complementary to the target sequences to hybridize under appropriate reaction conditions, particularly high stringency conditions, as outlined herein.

A nucleic acid probe is generally single stranded but can be partially single and partially double stranded. The strandedness of the probe is dictated by the structure, composition, and properties of the target sequence. In general, the nucleic acid probes range from about 8 to about 100 bases long, with from about 10 to about 80 bases being preferred, and from about 30 to about 50 bases being particularly preferred. That is, generally complements of ORFs or whole genes are not used. In some embodiments, nucleic acids of lengths up to hundreds of bases can be used.

In a preferred embodiment, more than one probe per sequence is used, with either overlapping probes or probes to different sections of the target being used. That is, two, three, four or more probes, with three being preferred, are used to build in a redundancy for a

particular target. The probes can be overlapping (i.e., have some sequence in common), or separate. In some cases, PCR primers may be used to amplify signal for higher sensitivity.

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As will be appreciated by those in the art, nucleic acids can be attached or immobilized to a solid support in a wide variety of ways. By "immobilized" and grammatical equivalents herein is meant the association or binding between the nucleic acid probe and the solid support is sufficient to be stable under the conditions of binding, washing, analysis, and removal as outlined below. The binding can typically be covalent or non-covalent. By "non-covalent binding" and grammatical equivalents herein is typically meant one or more of electrostatic, hydrophilic, and hydrophobic interactions. Included in non-covalent binding is the covalent attachment of a molecule, such as, streptavidin to the support and the non-covalent binding of the biotinylated probe to the streptavidin. By "covalent binding" and grammatical equivalents herein is meant that the two moieties, the solid support and the probe, are attached by at least one bond, including sigma bonds, pi bonds and coordination bonds. Covalent bonds can be formed directly between the probe and the solid support or can be formed by a cross linker or by inclusion of a specific reactive group on either the solid support or the probe or both molecules. Immobilization may also involve a combination of covalent and non-covalent interactions.

In general, the probes are attached to a biochip in a wide variety of ways, as will be appreciated by those in the art. As described herein, the nucleic acids can either be synthesized first, with subsequent attachment to the biochip, or can be directly synthesized on the biochip.

The biochip comprises a suitable solid substrate. By "substrate" or "solid support" or other grammatical equivalents herein is meant a material that can be modified for the attachment or association of the nucleic acid probes and is amenable to at least one detection method. Often the substrate may contain discrete individual sites appropriate for ndivitual partitioning and identification. As will be appreciated by those in the art, the number of possible substrates are very large, and include, but are not limited to, glass and modified or functionalized glass, plastics (including acrylics, polystyrene and copolymers of styrene and other materials, polypropylene, polyethylene, polybutylene, polyurethanes, Teflon, etc.), polysaccharides, nylon or nitrocellulose, resins, silica or silica-based materials including silicon and modified silicon, carbon, metals, inorganic glasses, plastics, etc. In general, the substrates allow optical detection and do not appreciably fluoresce. A preferred substrate is described in US application entitled Reusable Low Fluorescent Plastic Biochip, U.S.

WO 02/086443 PCT/US02/12476 Application Serial No. 09/270,214, filed March 15, 1999, herein incorporated by reference in its entirety.

Generally the substrate is planar, although as will be appreciated by those in the art, other configurations of substrates may be used as well. For example, the probes may be placed on the inside surface of a tube, for flow-through sample analysis to minimize sample volume. Similarly, the substrate may be flexible, such as a flexible foam, including closed cell foams made of particular plastics.

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In a preferred embodiment, the surface of the biochip and the probe may be derivatized with chemical functional groups for subsequent attachment of the two. Thus, e.g., the biochip is derivatized with a chemical functional group including, but not limited to, amino groups, carboxy groups, oxo groups and thiol groups, with amino groups being particularly preferred. Using these functional groups, the probes can be attached using functional groups on the probes. For example, nucleic acids containing amino groups can be attached to surfaces comprising amino groups, e.g., using linkers as are known in the art; e.g., homo-or hetero-bifunctional linkers as are well known (see 1994 Pierce Chemical Company catalog, technical section on cross-linkers, pages 155-200). In addition, in some cases, additional linkers, such as alkyl groups (including substituted and heteroalkyl groups) may be used.

In this embodiment, oligonucleotides are synthesized, and then attached to the surface of the solid support. Either the 5' or 3' terminus may be attached to the solid support, or attachment may be via linkage to an internal nucleoside.

In another embodiment, the immobilization to the solid support may be very strong, yet non-covalent. For example, biotinylated oligonucleotides can be made, which bind to surfaces covalently coated with streptavidin, resulting in attachment.

Alternatively, the oligonucleotides may be synthesized on the surface, as is known in the art. For example, photoactivation techniques utilizing photopolymerization compounds and techniques are used. In a preferred embodiment, the nucleic acids can be synthesized *in situ*, using known photolithographic techniques, such as those described in WO 95/25116; WO 95/35505; U.S. Patent Nos. 5,700,637 and 5,445,934; and references cited within, all of which are expressly incorporated by reference; these methods of attachment form the basis of the Affymetrix GeneChipTM technology.

Often, amplification-based assays are performed to measure the expression level of lung cancer-associated sequences. These assays are typically performed in conjunction with

reverse transcription. In such assays, a lung cancer-associated nucleic acid sequence acts as a template in an amplification reaction (e.g., Polymerase Chain Reaction, or PCR). In a quantitative amplification, the amount of amplification product will be proportional to the amount of template in the original sample. Comparison to appropriate controls provides a measure of the amount of lung cancer-associated RNA. Methods of quantitative amplification are well known to those of skill in the art. Detailed protocols for quantitative PCR are provided, e.g., in Innis, et al. (1990) PCR Protocols, A Guide to Methods and Applications.

In some embodiments, a TaqMan based assay is used to measure expression. TaqMan based assays use a fluorogenic oligonucleotide probe that contains a 5' fluorescent dye and a 3' quenching agent. The probe hybridizes to a PCR product, but cannot itself be extended due to a blocking agent at the 3' end. When the PCR product is amplified in subsequent cycles, the 5' nuclease activity of the polymerase, e.g., AmpliTaq, results in the cleavage of the TaqMan probe. This cleavage separates the 5' fluorescent dye and the 3' quenching agent, thereby resulting in an increase in fluorescence as a function of amplification (see, e.g., literature provided by Perkin-Elmer, e.g., www2.perkin-elmer.com).

Other suitable amplification methods include, but are not limited to, ligase chain reaction (LCR) (see Wu and Wallace (1989) Genomics 4:560, Landegren, et al. (1988) Science 241:1077, and Barringer, et al. (1990) Gene 89:117), transcription amplification (Kwoh, et al. (1989) Proc. Natl. Acad. Sci. USA 86:1173), self-sustained sequence replication (Guatelli, et al. (1990) Proc. Nat. Acad. Sci. USA 87:1874), dot PCR, and linker adapter PCR, etc.

Expression of lung cancer proteins from nucleic acids

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In a preferred embodiment, lung cancer nucleic acids, e.g., encoding lung cancer proteins, are used to make a variety of expression vectors to express lung cancer proteins which can then be used in screening assays, as described below. Expression vectors and recombinant DNA technology are well known to those of skill in the art (see, e.g., Ausubel, supra, and Fernandez and Hoeffler (eds 1999) Gene Expression Systems) and are used to express proteins. The expression vectors may be either self-replicating extrachromosomal vectors or vectors which integrate into a host genome. Generally, these expression vectors include transcriptional and translational regulatory nucleic acid operably linked to the nucleic acid encoding the lung cancer protein. The term "control sequences" refers to DNA

sequences used for the expression of an operably linked coding sequence in a particular host organism. Control sequences that are suitable for prokaryotes, e.g., include a promoter, optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

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Nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous, and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is typically accomplished by ligation at convenient restriction sites. If such sites do not exist, synthetic oligonucleotide adaptors or linkers are used in accordance with conventional practice. Transcriptional and translational regulatory nucleic acid will generally be appropriate to the host cell used to express the lung cancer protein. Numerous types of appropriate expression vectors, and suitable regulatory sequences are known in the art for a variety of host cells.

In general, transcriptional and translational regulatory sequences may include, but are not limited to, promoter sequences, ribosomal binding sites, transcriptional start and stop sequences, translational start and stop sequences, and enhancer or activator sequences. In a preferred embodiment, the regulatory sequences include a promoter and transcriptional start and stop sequences.

Promoter sequences may be either constitutive or inducible promoters. The promoters may be either naturally occurring promoters or hybrid promoters. Hybrid promoters, which combine elements of more than one promoter, are also known in the art, and are useful in the present invention.

In addition, an expression vector may comprise additional elements. For example, the expression vector may have two replication systems, thus allowing it to be maintained in two organisms, e.g., in mammalian or insect cells for expression and in a prokaryotic host for cloning and amplification. Furthermore, for integrating expression vectors, the expression vector often contains at least one sequence homologous to the host cell genome, and preferably two homologous sequences which flank the expression construct. The integrating

vector may be directed to a specific locus in the host cell by selecting the appropriate homologous sequence for inclusion in the vector. Constructs for integrating vectors are well known in the art (e.g., Fernandez and Hoeffler, *supra*).

In addition, in a preferred embodiment, the expression vector contains a selectable marker gene to allow the selection of transformed host cells. Selection genes are well known in the art and will vary with the host cell used.

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The lung cancer proteins of the present invention are usually produced by culturing a host cell transformed with an expression vector containing nucleic acid encoding a lung cancer protein, under the appropriate conditions to induce or cause expression of the lung cancer protein. Conditions appropriate for lung cancer protein expression will vary with the choice of the expression vector and the host cell, and will be easily ascertained by one skilled in the art through routine experimentation or optimization. For example, the use of constitutive promoters in the expression vector will require optimizing the growth and proliferation of the host cell, while the use of an inducible promoter requires the appropriate growth conditions for induction. In addition, in some embodiments, the timing of the harvest is important. For example, the baculoviral systems used in insect cell expression are lytic viruses, and thus harvest time selection can be crucial for product yield.

Appropriate host cells include yeast, bacteria, archaebacteria, fungi, and insect and animal cells, including mammalian cells. Of particular interest are *Saccharomyces cerevisiae* and other yeasts, *E. coli*, *Bacillus subtilis*, Sf9 cells, C129 cells, 293 cells, *Neurospora*, BHK, CHO, COS, HeLa cells, HUVEC (human umbilical vein endothelial cells), THP1 cells (a macrophage cell line) and various other human cells and cell lines.

In a preferred embodiment, the lung cancer proteins are expressed in mammalian cells. Mammalian expression systems are also known in the art, and include retroviral and adenoviral systems. Of particular use as mammalian promoters are the promoters from mammalian viral genes, since the viral genes are often highly expressed and have a broad host range. Examples include the SV40 early promoter, mouse mammary tumor virus LTR promoter, adenovirus major late promoter, herpes simplex virus promoter, and the CMV promoter (see, e.g., Fernandez and Hoeffler, *supra*). Typically, transcription termination and polyadenylation sequences recognized by mammalian cells are regulatory regions located 3' to the translation stop codon and thus, together with the promoter elements, flank the coding sequence. Examples of transcription terminator and polyadenylation signals include those derived form SV40.

The methods of introducing exogenous nucleic acid into mammalian hosts, as well as other hosts, is well known in the art, and will vary with the host cell used. Techniques include dextran-mediated transfection, calcium phosphate precipitation, polybrene mediated transfection, protoplast fusion, electroporation, viral infection, encapsulation of the polynucleotide(s) in liposomes, and direct microinjection of the DNA into nuclei.

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In a preferred embodiment, lung cancer proteins are expressed in bacterial systems. Promoters from bacteriophage may also be used and are known in the art. In addition, synthetic promoters and hybrid promoters are also useful; e.g., the tac promoter is a hybrid of the trp and lac promoter sequences. Furthermore, a bacterial promoter can include naturally occurring promoters of non-bacterial origin that have the ability to bind bacterial RNA polymerase and initiate transcription. In addition to a functioning promoter sequence, an efficient ribosome binding site is desirable. The expression vector may also include a signal peptide sequence that provides for secretion of the lung cancer protein in bacteria. The protein is either secreted into the growth media (gram-positive bacteria) or into the periplasmic space, located between the inner and outer membrane of the cell (gram-negative bacteria). The bacterial expression vector may also include a selectable marker gene to allow for the selection of bacterial strains that have been transformed. Suitable selection genes include genes which render the bacteria resistant to drugs such as ampicillin, chloramphenicol, erythromycin, kanamycin, neomycin and tetracycline. Selectable markers also include biosynthetic genes, such as those in the histidine, tryptophan and leucine biosynthetic pathways. These components are assembled into expression vectors. Expression vectors for bacteria are well known in the art, and include vectors for Bacillus subtilis, E. coli, Streptococcus cremoris, and Streptococcus lividans, among others (e.g., Fernandez and Hoeffler, supra). The bacterial expression vectors are transformed into bacterial host cells using techniques well known in the art, such as calcium chloride treatment, electroporation, and others.

In one embodiment, lung cancer proteins are produced in insect cells. Expression vectors for the transformation of insect cells, and in particular, baculovirus-based expression vectors, are well known in the art.

In a preferred embodiment, lung cancer protein is produced in yeast cells. Yeast expression systems are well known in the art, and include expression vectors for Saccharomyces cerevisiae, Candida albicans and C. maltosa, Hansenula polymorpha,

Kluyveromyces fragilis and K. lactis, Pichia guillerimondii, and P. pastoris, Schizosaccharomyces pombe, and Yarrowia lipolytica.

The lung cancer protein may also be made as a fusion protein, using techniques well known in the art. Thus, e.g., for the creation of monoclonal antibodies, if the desired epitope is small, the lung cancer protein may be fused to a carrier protein to form an immunogen. Alternatively, the lung cancer protein may be made as a fusion protein to increase expression for affinity purification purposes, or for other reasons. For example, when the lung cancer protein is a lung cancer peptide, the nucleic acid encoding the peptide may be linked to other nucleic acid for expression purposes.

In a preferred embodiment, the lung cancer protein is purified or isolated after expression. Lung cancer proteins may be isolated or purified in a variety of appropriate ways. Standard purification methods include electrophoretic, molecular, immunological and chromatographic techniques, including ion exchange, hydrophobic, affinity, and reverse-phase HPLC chromatography, and chromatofocusing. For example, the lung cancer protein may be purified using a standard anti-lung cancer protein antibody column. Ultrafiltration and diafiltration techniques, in conjunction with protein concentration, are also useful. For general guidance in suitable purification techniques, see Scopes (1982) Protein Purification. The degree of purification necessary will vary depending on the use of the lung cancer protein. In some instances no purification will be necessary.

Once expressed and purified if necessary, the lung cancer proteins and nucleic acids are useful in a number of applications. They may be used as immunoselection reagents, as vaccine reagents, as screening agents, therapeutic entities, for production of antibodies, as transcription or translation inhibitors, etc.

25 Variants of lung cancer proteins

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In one embodiment, the lung cancer proteins are derivative or variant lung cancer proteins as compared to the wild-type sequence. That is, as outlined more fully below, the derivative lung cancer peptide will often contain at least one amino acid substitution, deletion or insertion, with amino acid substitutions being particularly preferred. The amino acid substitution, insertion or deletion may occur at a particular residue within the lung cancer peptide.

Also included within one embodiment of lung cancer proteins of the present invention are amino acid sequence variants. These variants typically fall into one or more of three

WO 02/086443 PCT/US02/12476 classes: substitutional, insertional or deletional variants. These variants ordinarily are prepared by site specific mutagenesis of nucleotides in the DNA encoding the lung cancer protein, using cassette or PCR mutagenesis or other techniques, to produce DNA encoding the variant, and thereafter expressing the DNA in recombinant cell culture as outlined above. However, variant lung cancer protein fragments having up to about 100-150 residues may be

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although variants can also be selected which have modified characteristics as will be more fully outlined below.

prepared by in vitro synthesis. Amino acid sequence variants are characterized by the

predetermined nature of the variation, a feature that sets them apart from naturally occurring

allelic or interspecies variation of the lung cancer protein amino acid sequence. The variants

typically exhibit a similar qualitative biological activity as the naturally occurring analogue,

While the site or region for introducing an amino acid sequence variation is often predetermined, the mutation per se need not be predetermined. For example, in order to optimize the performance of a mutation at a given site, random mutagenesis may be conducted at the target codon or region and the expressed lung cancer variants screened for the optimal combination of desired activity. Techniques exist for making substitution mutations at predetermined sites in DNA having a known sequence, e.g., M13 primer mutagenesis and PCR mutagenesis. Screening of mutants is often done using assays of lung cancer protein activities.

Amino acid substitutions are typically of single residues; insertions usually will be on the order of from about 1 to 20 amino acids, although considerably larger insertions may be occasionally tolerated. Deletions generally range from about 1 to about 20 residues, although in some cases deletions may be much larger.

Substitutions, deletions, insertions or any combination thereof may be used to arrive at a final derivative. Generally these changes are done on a few amino acids to minimize the alteration of the molecule. Larger changes may be tolerated in certain circumstances. When small alterations in the characteristics of a lung cancer protein are desired, substitutions are generally made in accordance with the amino acid substitution chart provided in the definition section.

Variants typically exhibit essentially the same qualitative biological activity and will elicit the same immune response as a naturally-occurring analog, although variants also are selected to modify the characteristics of lung cancer proteins as needed. Alternatively, the

variant may be designed or reorganized such that a biological activity of the lung cancer protein is altered. For example, glycosylation sites may be added, altered, or removed.

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Covalent modifications of lung cancer polypeptides are included within the scope of this invention. One type of covalent modification includes reacting targeted amino acid residues of a lung cancer polypeptide with an organic derivatizing agent that is capable of reacting with selected side chains or the N-or C-terminal residues of a lung cancer polypeptide. Derivatization with bifunctional agents is useful, for instance, for crosslinking lung cancer polypeptides to a water-insoluble support matrix or surface for use in a method for purifying anti-lung cancer polypeptide antibodies or screening assays, as is more fully described below. Commonly used crosslinking agents include, e.g., 1,1-bis(diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, e.g., esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), bifunctional maleimides such as bis-N-maleimido-1,8-octane and agents such as methyl-3-((p-azidophenyl)dithio)propioimidate.

Other modifications include deamidation of glutaminyl and asparaginyl residues to the corresponding glutamyl and aspartyl residues, respectively, hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of serinyl, threonyl or tyrosyl residues, methylation of the γ -amino groups of lysine, arginine, and histidine side chains (Creighton (1983) Proteins: Structure and Molecular Properties, pp. 79-86), acetylation of the N-terminal amine, and amidation of any C-terminal carboxyl group.

Another type of covalent modification of the lung cancer polypeptide encompassed by this invention is an altered native glycosylation pattern of the polypeptide. "Altering the native glycosylation pattern" is intended herein to mean adding to or deleting one or more carbohydrate moieties of a native sequence lung cancer polypeptide. Glycosylation patterns can be altered in many ways. For example the use of different cell types to express lung cancer-associated sequences can result in different glycosylation patterns.

Addition of glycosylation sites to lung cancer polypeptides may also be accomplished by altering the amino acid sequence thereof. The alteration may be made, e.g., by the addition of, or substitution by, one or more serine or threonine residues to the native sequence lung cancer polypeptide (for O-linked glycosylation sites). The lung cancer amino acid sequence may optionally be altered through changes at the DNA level, particularly by mutating the DNA encoding the lung cancer polypeptide at preselected bases such that codons are generated that will translate into the desired amino acids.

Another means of increasing the number of carbohydrate moieties on the lung cancer polypeptide is by chemical or enzymatic coupling of glycosides to the polypeptide. Such methods are described in the art, e.g., in WO 87/05330, and in Aplin and Wriston (1981) . CRC Crit. Rev. Biochem., pp. 259-306.

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Removal of carbohydrate moieties present on the lung cancer polypeptide may be accomplished chemically or enzymatically or by mutational substitution of codons encoding for amino acid residues that serve as targets for glycosylation. Chemical deglycosylation techniques are known in the art and described, for instance, by Hakimuddin, et al. (1987)

Arch. Biochem. Biophys., 259:52 and by Edge, et al. (1981) Anal. Biochem., 118:131.

Enzymatic cleavage of carbohydrate moieties on polypeptides can be achieved by the use of a variety of endo-and exo-glycosidases as described by Thotakura, et al. (1987) Meth.

Enzymol., 138:350.

Another type of covalent modification of lung cancer comprises linking the lung cancer polypeptide to one of a variety of nonproteinaceous polymers, e.g., polyethylene glycol, polypropylene glycol, or polyoxyalkylenes, in the manner set forth in U.S. Patent Nos. 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192, or 4,179,337.

Lung cancer polypeptides of the present invention may also be modified in a way to form chimeric molecules comprising a lung cancer polypeptide fused to another, heterologous polypeptide or amino acid sequence. In one embodiment, such a chimeric molecule comprises a fusion of a lung cancer polypeptide with a tag polypeptide which provides an epitope to which an anti-tag antibody can selectively bind. The epitope tag is generally placed at the amino-or carboxyl-terminus of the lung cancer polypeptide. The presence of such epitope-tagged forms of a lung cancer polypeptide can be detected using an antibody against the tag polypeptide. Also, provision of the epitope tag enables the lung cancer polypeptide to be readily purified by affinity purification using an anti-tag antibody or another type of affinity matrix that binds to the epitope tag. In an alternative embodiment, the chimeric molecule may comprise a fusion of a lung cancer polypeptide with an immunoglobulin or a particular region of an immunoglobulin. For a bivalent form of the chimeric molecule, such a fusion could be to the Fc region of an IgG molecule.

Various tag polypeptides and their respective antibodies are well known and examples include poly-histidine (poly-his) or poly-histidine-glycine (poly-his-gly) tags; HIS6 and metal chelation tags, the flu HA tag polypeptide and its antibody 12CA5 (Field, et al. (1988) Mol. Cell. Biol. 8:2159-2165); the c-myc tag and the 8F9, 3C7, 6E10, G4, B7 and 9E10 antibodies

thereto (Evan, et al. (1985) Molecular and Cellular Biology 5:3610-3616); and the Herpes Simplex virus glycoprotein D (gD) tag and its antibody (Paborsky, et al. (1990) Protein Engineering 3(6):547-553). Other tag polypeptides include the Flag-peptide (Hopp, et al. (1988) BioTechnology 6:1204-1210); the KT3 epitope peptide (Martin, et al. (1992) Science 255:192-194); tubulin epitope peptide (Skinner, et al. (1991) J. Biol. Chem. 266:15163-15166); and the T7 gene 10 protein peptide tag (Lutz-Freyermuth, et al. (1990) Proc. Nat'l

Also included are other lung cancer proteins of the lung cancer family, and lung cancer proteins from other organisms, which are cloned and expressed as outlined below. Thus, probe or degenerate polymerase chain reaction (PCR) primer sequences may be used to find other related lung cancer proteins from primates or other organisms. As will be appreciated by those in the art, particularly useful probe and/or PCR primer sequences include unique areas of the lung cancer nucleic acid sequence. As is generally known in the art, preferred PCR primers are from about 15 to about 35 nucleotides in length, with from about 20 to about 30 being preferred, and may contain inosine as needed. PCR reaction conditions are well known in the art (e.g., Innis, PCR Protocols, supra).

Antibodies to lung cancer proteins

Acad. Sci. USA 87:6393-6397).

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In a preferred embodiment, when a lung cancer protein is to be used to generate antibodies, e.g., for immunotherapy or immunodiagnosis, the lung cancer protein should share at least one epitope or determinant with the full length protein. By "epitope" or "determinant" herein is typically meant a portion of a protein which will generate and/or bind an antibody or T-cell receptor in the context of MHC. Thus, in most instances, antibodies made to a smaller lung cancer protein will be able to bind to the full-length protein, particularly linear epitopes. In a preferred embodiment, the epitope is unique; that is, antibodies generated to a unique epitope show little or no cross-reactivity.

Methods of preparing polyclonal antibodies are well known (e.g., Coligan, supra; and Harlow and Lane, supra). Polyclonal antibodies can be raised in a mammal, e.g., by one or more injections of an immunizing agent and, if desired, an adjuvant. Typically, the immunizing agent and/or adjuvant will be injected in the mammal by multiple subcutaneous or intraperitoneal injections. The immunizing agent may include a protein encoded by a nucleic acid of Tables 1A-16 or fragment thereof or a fusion protein thereof. It may be useful to conjugate the immunizing agent to a protein known to be immunogenic in the mammal

being immunized. Immunogenic proteins include, e.g., keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. Adjuvants include, e.g., Freund's complete adjuvant and MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate). The immunization protocol may be selected by one skilled in the art.

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The antibodies may, alternatively, be monoclonal antibodies. Monoclonal antibodies may be prepared using hybridoma methods, such as those described by Kohler and Milstein (1975) Nature 256:495. In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes may be immunized in vitro. The immunizing agent will typically include a polypeptide encoded by a nucleic acid of the tables, or fragment thereof, or a fusion protein thereof. Generally, either peripheral blood lymphocytes ("PBLs") are used if cells of human origin are desired, or spleen cells or lymph node cells are used if nonhuman mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding (1986) Monoclonal Antibodies: Principles and Practice, pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovin, or primate origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells may be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

In one embodiment, the antibodies are bispecific antibodies. Bispecific antibodies are typically monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens or that have binding specificities for two epitopes on the same antigen. In one embodiment, one of the binding specificities is for a protein encoded by a nucleic acid of the tables or a fragment thereof, the other one is for any other antigen, and preferably for a cell-surface protein or receptor or receptor subunit, preferably one that is tumor specific. Alternatively, tetramer-type technology may create multivalent reagents.

In a preferred embodiment, the antibodies to lung cancer protein are capable of reducing or eliminating a biological function of a lung cancer protein, in a naked form or conjugated to an effector moiety. That is, the addition of anti-lung cancer protein antibodies (either polyclonal or preferably monoclonal) to lung cancer tissue (or cells containing lung cancer) may reduce or eliminate the lung cancer. Generally, at least a 25% decrease in activity, growth, size or the like is preferred, with at least about 50% being particularly preferred and about a 95-100% decrease being especially preferred.

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In a preferred embodiment the antibodies to the lung cancer proteins are humanized antibodies (e.g., Xenerex Biosciences, Medarex, Inc., Abgenix, Inc., Protein Design Labs, Inc.) Humanized forms of non-human (e.g., murine) antibodies are chimeric molecules of immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')2 or other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. Humanized antibodies include human immunoglobulins (recipient antibody) in which residues from a complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat or rabbit having the desired specificity, affinity and capacity. In some instances, Fv framework residues of a human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies may also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, a humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework (FR) regions are those of a human immunoglobulin consensus sequence. A humanized antibody optimally also will typically comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones, et al. (1986) Nature 321:522-525; Riechmann, et al. (1988) Nature 332:323-329; and Presta (1992) Curr. Op. Struct. Biol. 2:593-596). Humanization can be performed following the method of Winter and co-workers (Jones, et al. (1986) Nature 321:522-525; Riechmann, et al. (1988) Nature 332:323-327; Verhoeyen, et al. (1988) Science 239:1534-1536), by substituting rodent CDRs or CDR sequences for corresponding sequences of a human antibody. Accordingly, such humanized antibodies are chimeric antibodies (U.S. Patent No. 4,816,567), wherein substantially less than an intact human variable domain has been substituted by corresponding sequence from a non-human species.

Human-like antibodies can also be produced using various techniques known in the art, including phage display libraries (Hoogenboom and Winter (1991) J. Mol. Biol. 227:381; Marks, et al. (1991) J. Mol. Biol. 222:581). The techniques of Cole, et al. and Boerner, et al. are also available for the preparation of human monoclonal antibodies (Cole, et al. (1985) 5 Monoclonal Antibodies and Cancer Therapy, p. 77 and Boerner, et al. (1991) J. Immunol. 147(1):86-95). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in nearly all respects, including gene rearrangement, assembly, and antibody repertoire. This 10 approach is described, e.g., in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in the following scientific publications: Marks, et al. (1992) Bio/Technology 10:779-783; Lonberg, et al. (1994) Nature 368:856-859; Morrison (1994) Nature 368:812-13; Fishwild, et al. (1996) Nature Biotechnology 14:845-51; Neuberger (1996) Nature Biotechnology 14:826; and Lonberg and Huszar (1995) Intern. Rev. Immunol. 15 13:65-93.

By immunotherapy is meant treatment of lung cancer with an antibody raised against a lung cancer proteins. As used herein, immunotherapy can be passive or active. Passive immunotherapy as defined herein is the passive transfer of antibody to a recipient (patient). Active immunization is the induction of antibody and/or T-cell responses in a recipient (patient). Induction of an immune response is the result of providing the recipient with an antigen to which antibodies are raised. The antigen may be provided by injecting a polypeptide against which antibodies are desired to be raised into a recipient, or contacting the recipient with a nucleic acid capable of expressing the antigen and under conditions for expression of the antigen, leading to an immune response.

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In a preferred embodiment the lung cancer proteins against which antibodies are raised are secreted proteins as described above. Without being bound by theory, antibodies used for treatment, may bind and prevent the secreted protein from binding to its receptor, thereby inactivating the secreted lung cancer protein.

In another preferred embodiment, the lung cancer protein to which antibodies are raised is a transmembrane protein. Without being bound by theory, antibodies used for treatment may bind the extracellular domain of the lung cancer protein and prevent it from binding to other proteins, such as circulating ligands or cell-associated molecules. The

antibody may cause down-regulation of the transmembrane lung cancer protein. The antibody may be a competitive, non-competitive or uncompetitive inhibitor of protein binding to the extracellular domain of the lung cancer protein. The antibody may be an antagonist of the lung cancer protein or may prevent activation of a transmembrane lung cancer protein, or may induce or suppress a particular cellular pathway. In some embodiments, when the antibody prevents the binding of other molecules to the lung cancer protein, the antibody prevents growth of the cell. The antibody may also be used to target or sensitize the cell to cytotoxic agents, including, but not limited to TNF-α, TNF-β, IL-1, INF-γ, and IL-2, or chemotherapeutic agents including 5FU, vinblastine, actinomycin D, cisplatin, methotrexate, and the like. In some instances the antibody may belong to a sub-type that activates serum complement when complexed with the transmembrane protein thereby mediating cytotoxicity or antigen-dependent cytotoxicity (ADCC). Thus, lung cancer may be treated by administering to a patient antibodies directed against the transmembrane lung cancer protein. Antibody-labeling may activate a co-toxin, localize a toxin payload, or otherwise provide means to locally ablate cells.

In another preferred embodiment, the antibody is conjugated to an effector moiety. The effector moiety can be various molecules, including labeling moieties such as radioactive labels or fluorescent labels, or can be a therapeutic moiety. In one aspect the therapeutic moiety is a small molecule that modulates the activity of a lung cancer protein. In another aspect the therapeutic moiety may modulate an activity of molecules associated with or in close proximity to a lung cancer protein. The therapeutic moiety may inhibit enzymatic or signaling activity such as protease or collagenase activity associated with lung cancer.

In a preferred embodiment, the therapeutic moiety can also be a cytotoxic agent. In this method, targeting the cytotoxic agent to lung cancer tissue or cells results in a reduction in the number of afflicted cells, thereby reducing symptoms associated with lung cancer. Cytotoxic agents are numerous and varied and include, but are not limited to, cytotoxic drugs or toxins or active fragments of such toxins. Suitable toxins and their corresponding fragments include diphtheria A chain, exotoxin A chain, ricin A chain, abrin A chain, curcin, crotin, phenomycin, enomycin, saporin, auristatin, and the like. Cytotoxic agents also include radiochemicals made by conjugating radioisotopes to antibodies raised against lung cancer proteins, or binding of a radionuclide to a chelating agent that has been covalently attached to the antibody. Targeting the therapeutic moiety to transmembrane lung cancer proteins not only serves to increase the local concentration of therapeutic moiety in the lung cancer

WO 02/086443 PCT/US02/12476 afflicted area, but also serves to reduce deleterious side effects that may be associated with the untargeted therapeutic moiety.

In another preferred embodiment, the lung cancer protein against which the antibodies are raised is an intracellular protein. In this case, the antibody may be conjugated to a protein or other entity which facilitates entry into the cell. In one case, the antibody enters the cell by endocytosis. In another embodiment, a nucleic acid encoding the antibody is administered to the individual or cell. Moreover, wherein the lung cancer protein can be targeted within a cell, i.e., the nucleus, an antibody theretomay contain a signal for that target localization, i.e., a nuclear localization signal.

The lung cancer antibodies of the invention specifically bind to lung cancer proteins. By "specifically bind" herein is meant that the antibodies bind to the protein with a K_d of at least about 0.1 mM, more usually at least about 1 µM, preferably at least about 0.1 µM or better, and most preferably, 0.01 µM or better. Selectivity of binding to the specific target and not to related other sequences is also important.

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Detection of lung cancer sequence for diagnostic and therapeutic applications

In one aspect, the RNA expression levels of genes are determined for different cellular states in the lung cancer phenotype. Expression levels of genes in normal tissue (e.g., not undergoing lung cancer), in lung cancer tissue (and in some cases, for varying severities of lung cancer that relate to prognosis, as outlined below), or in non-malignant disease are evaluated to provide expression profiles. A gene expression profile of a particular cell state or point of development is essentially a "fingerprint" of the state of the cell. While two states may have a particular gene similarly expressed, the evaluation of a number of genes simultaneously allows the generation of a gene expression profile that is reflective of the state of the cell. By comparing expression profiles of cells in different states, information regarding which genes are important (including both up- and down-regulation of genes) in each of these states is obtained. Then, diagnosis may be performed or confirmed to determine whether a tissue sample has the gene expression profile of normal or cancerous tissue. This will provide for molecular diagnosis of related conditions.

"Differential expression," or grammatical equivalents as used herein, refers to qualitative or quantitative differences in the temporal and/or cellular gene expression patterns within and among cells and tissue. Thus, a differentially expressed gene can qualitatively have its expression altered, including an activation or inactivation, in, e.g.,

WO 02/086443 PCT/US02/12476 normal versus lung cancer tissue. Genes may be turned on or turned off in a particular state, relative to another state thus permitting comparison of two or more states. A qualitatively regulated gene will exhibit an expression pattern within a state or cell type which is detectable by standard techniques. Some genes will be expressed in one state or cell type, but not in both. Alternatively, the difference in expression may be quantitative, e.g., in that expression is increased or decreased; i.e., gene expression is either upregulated, resulting in an increased amount of transcript, or downregulated, resulting in a decreased amount of transcript. The degree to which expression differs need only be large enough to quantify via standard characterization techniques as outlined below, such as by use of Affymetrix GeneChip™ expression arrays, Lockhart (1996) Nature Biotechnology 14:1675-1680, hereby expressly incorporated by reference. Other techniques include, but are not limited to, quantitative reverse transcriptase PCR, northern analysis and RNase protection. As outlined above, preferably the change in expression (i.e., upregulation or downregulation) is typically at least about 50%, more preferably at least about 100%, more preferably at least about 150%, more preferably at least about 200%, with from 300 to at least 1000% being especially preferred.

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Evaluation may be at the gene transcript or the protein level. The amount of gene expression may be monitored using nucleic acid probes to the RNA or DNA equivalent of the gene transcript, and the quantification of gene expression levels, or, alternatively, the final gene product itself (protein) can be monitored, e.g., with antibodies to the lung cancer protein and standard immunoassays (ELISAs, etc.) or other techniques, including mass spectroscopy assays, 2D gel electrophoresis assays, etc. Proteins corresponding to lung cancer genes, e.g., those identified as being important in a lung cancer or disease phenotype, can be evaluated in a lung cancer diagnostic test. In a preferred embodiment, gene expression monitoring is performed simultaneously on a number of genes.

The lung cancer nucleic acid probes may be attached to biochips as outlined herein for the detection and quantification of lung cancer sequences in a particular cell. The assays are further described below in the example. PCR techniques can be used to provide greater sensitivity. Multiple protein expression monitoring can be performed as well. Similarly, these assays may be performed on an individual basis as well.

In a preferred embodiment nucleic acids encoding the lung cancer protein are detected. Although DNA or RNA encoding the lung cancer protein may be detected, of particular interest are methods wherein an mRNA encoding a lung cancer protein is detected.

Probes to detect mRNA can be a nucleotide/deoxynucleotide probe that is complementary to and hybridizes with the mRNA and includes, but is not limited to, oligonucleotides, cDNA or RNA. Probes also should contain a detectable label, as defined herein. In one method the mRNA is detected after immobilizing the nucleic acid to be examined on a solid support such as nylon membranes and hybridizing the probe with the sample. Following washing to remove the non-specifically bound probe, the label is detected. In another method detection of the mRNA is performed *in situ*. In this method permeabilized cells or tissue samples are contacted with a detectably labeled nucleic acid probe for sufficient time to allow the probe to hybridize with the target mRNA. Following washing to remove the non-specifically bound probe, the label is detected. For example a digoxygenin labeled riboprobe (RNA probe) that is complementary to the mRNA encoding a lung cancer protein is detected by binding the digoxygenin with an anti-digoxygenin secondary antibody and developed with nitro blue tetrazolium and 5-bromo-4-chloro-3-indoyl phosphate.

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In a preferred embodiment, various proteins from the three classes of proteins as described herein (secreted, transmembrane or intracellular proteins) are used in diagnostic assays. The lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in diagnostic assays. This can be performed on an individual gene or corresponding polypeptide level. In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes and/or corresponding polypeptides.

As described and defined herein, lung cancer proteins, including intracellular, transmembrane, or secreted proteins, find use as markers of lung cancer, e.g., for prognostic or diagnostic purposes. Detection of these proteins in putative lung cancer tissue allows for detection, prognosis, or diagnosis of lung cancer or similar disease, and perhaps for selection of therapeutic strategy. In one embodiment, antibodies are used to detect lung cancer proteins. A preferred method separates proteins from a sample by electrophoresis on a gel (typically a denaturing and reducing protein gel, but may be another type of gel, including isoelectric focusing gels and the like). Following separation of proteins, the lung cancer protein is detected, e.g., by immunoblotting with antibodies raised against the lung cancer protein. Methods of immunoblotting are well known to those of ordinary skill in the art.

In another preferred method, antibodies to the lung cancer protein find use in *in situ* imaging techniques, e.g., in histology (e.g., Asai (ed. 1993) Methods in Cell Biology:

Antibodies in Cell Biology, volume 37. In this method cells are contacted with from one to many antibodies to the lung cancer protein(s). Following washing to remove non-specific antibody binding, the presence of the antibody or antibodies is detected. In one embodiment the antibody is detected by incubating with a secondary antibody that contains a detectable label, e.g., multicolor fluorescence or confocal imaging. In another method the primary antibody to the lung cancer protein(s) contains a detectable label, e.g., an enzyme marker that can act on a substrate. In another preferred embodiment each one of multiple primary antibodies contains a distinct and detectable label. This method finds particular use in simultaneous screening for a plurality of lung cancer proteins. Many other histological imaging techniques are also provided by the invention.

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In a preferred embodiment the label is detected in a fluorometer which has the ability to detect and distinguish emissions of different wavelengths. In addition, a fluorescence activated cell sorter (FACS) can be used in the method.

In another preferred embodiment, antibodies find use in diagnosing lung cancer from blood, serum, plasma, stool, and other samples. Such samples, therefore, are useful as samples to be probed or tested for the presence of lung cancer proteins. Antibodies can be used to detect a lung cancer protein by previously described immunoassay techniques including ELISA, immunoblotting (western blotting), immunoprecipitation, BIACORE technology and the like. Conversely, the presence of antibodies may indicate an immune response against an endogenous lung cancer protein or vaccine.

In a preferred embodiment, in situ hybridization of labeled lung cancer nucleic acid probes to tissue arrays is done. For example, arrays of tissue samples, including lung cancer tissue and/or normal tissue, are made. In situ hybridization (see, e.g., Ausubel, supra) is then performed. When comparing the fingerprints between an individual and a standard, the skilled artisan can make a diagnosis, a prognosis, or a prediction based on the findings. It is further understood that the genes which indicate the diagnosis may differ from those which indicate the prognosis and molecular profiling of the condition of the cells may lead to distinctions between responsive or refractory conditions or may be predictive of outcomes.

In a preferred embodiment, the lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in prognosis assays. As above, gene expression profiles can be generated that correlate to lung cancer, clinical, pathological, or other information, in terms of long term prognosis. Again, this may be done on either a protein or gene level, with the use of genes being preferred. Single or multiple

genes may be useful in various combinations. As above, lung cancer probes may be attached to biochips for the detection and quantification of lung cancer sequences in a tissue or patient. The assays proceed as outlined above for diagnosis. PCR method may provide more sensitive and accurate quantification.

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Assays for therapeutic compounds

candidate agent, see Zlokarnik, supra.

In a preferred embodiment, the proteins, nucleic acids, and antibodies as described herein are used in drug screening assays. The lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in drug screening assays or by evaluating the effect of drug candidates on a "gene expression profile" or expression profile of polypeptides. In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes after treatment with a candidate agent (e.g., Zlokarnik, et al. (1998) Science 279:84-8; Heid (1996) Genome Res. 6:986-94.

In a preferred embodiment, the lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing the native or modified lung cancer proteins are used in screening assays. That is, the present invention provides novel methods for screening for compositions which modulate the lung cancer phenotype or an identified physiological function of a lung cancer protein. As above, this can be done on an individual gene level or by evaluating the effect of drug candidates on a "gene expression profile". In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes after treatment with a

Having identified differentially expressed genes herein, a variety of assays may be performed. In a preferred embodiment, assays may be run on an individual gene or protein level. That is, having identified a particular gene with altered regulation in lung cancer, test compounds can be screened for the ability to modulate gene expression or for binding to the lung cancer protein. "Modulation" thus includes an increase or a decrease in gene expression. The preferred amount of modulation will depend on the original change of the gene expression in normal versus tissue undergoing lung cancer, with changes of at least 10%, preferably 50%, more preferably 100-300%, and in some embodiments 300-1000% or greater. Thus, if a gene exhibits a 4-fold increase in lung cancer tissue compared to normal tissue, a decrease of about four-fold is often desired; similarly, a 10-fold decrease in lung

cancer tissue compared to normal tissue often provides a target value of a 10-fold increase in expression to be induced by the test compound.

The amount of gene expression may be monitored using nucleic acid probes and the quantification of gene expression levels, or, alternatively, the gene product itself can be monitored, e.g., through the use of antibodies to the lung cancer protein and standard immunoassays. Proteomics and separation techniques may also allow quantification of expression.

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In a preferred embodiment, gene or protein expression monitoring of a number of entities, i.e., an expression profile, is monitored simultaneously. Such profiles will typically involve a plurality of those entities described herein.

In this embodiment, the lung cancer nucleic acid probes are attached to biochips as outlined herein for the detection and quantification of lung cancer sequences in a particular cell. Alternatively, PCR may be used. Thus, a series, e.g., of microtiter plate, may be used with dispensed primers in desired wells. A PCR reaction can then be performed and analyzed for each well.

Expression monitoring can be performed to identify compounds that modify the expression of one or more lung cancer-associated sequences, e.g., a polynucleotide sequence set out in the tables. Generally, in a preferred embodiment, a test compound is added to the cells prior to analysis. Moreover, screens are also provided to identify agents that modulate lung cancer, modulate lung cancer proteins, bind to a lung cancer protein, or interfere with the binding of a lung cancer protein and an antibody, substrate, or other binding partner.

The term "test compound" or "drug candidate" or "modulator" or grammatical equivalents as used herein describes a molecule, e.g., protein, oligopeptide, small organic molecule, polysaccharide, polynucleotide, etc., to be tested for the capacity to directly or indirectly alter the lung cancer phenotype or the expression of a lung cancer sequence, e.g., a nucleic acid or protein sequence. In preferred embodiments, modulators alter expression profiles of nucleic acids or proteins provided herein. In one embodiment, the modulator suppresses a lung cancer phenotype, e.g., to a normal or non-malignant tissue fingerprint. In another embodiment, a modulator induces a lung cancer phenotype. Generally, a plurality of assay mixtures are run in parallel with different agent concentrations to obtain a differential response to the various concentrations. Typically, one of these concentrations serves as a negative control, i.e., at zero concentration or below the level of detection.

In one aspect, a modulator will neutralize the effect of a lung cancer protein. By "neutralize" is meant that activity of a protein and the consequent effect on the cell is inhibited or blocked.

In certain embodiments, combinatorial libraries of potential modulators will be screened for an ability to bind to a lung cancer polypeptide or to modulate activity. Conventionally, new chemical entities with useful properties are generated by identifying a chemical compound (called a "lead compound") with some desirable property or activity, e.g., inhibiting activity, creating variants of the lead compound, and evaluating the property and activity of those variant compounds. Often, high throughput screening (HTS) methods are employed for such an analysis.

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In one preferred embodiment, high throughput screening methods involve providing a library containing a large number of potential therapeutic compounds (candidate compounds). Such "combinatorial chemical libraries" are then screened in one or more assays to identify those library members (particular chemical species or subclasses) that display a desired characteristic activity. The compounds thus identified can serve as conventional "lead compounds" or can themselves be used as potential or actual therapeutics.

A combinatorial chemical library is a collection of diverse chemical compounds generated by either chemical synthesis or biological synthesis by combining a number of chemical "building blocks" such as reagents. For example, a linear combinatorial chemical library, such as a polypeptide (e.g., mutein) library, is formed by combining a set of chemical building blocks called amino acids in every possible way for a given compound length (i.e., the number of amino acids in a polypeptide compound). Millions of chemical compounds can be synthesized through such combinatorial mixing of chemical building blocks (Gallop, et al. (1994) <u>J. Med. Chem.</u> 37(9):1233-1251).

Preparation and screening of combinatorial chemical libraries is well known to those of skill in the art. Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Patent No. 5,010,175, Furka (1991) Pept. Prot. Res. 37:487-493, Houghton, et al. (1991) Nature, 354:84-88), peptoids (PCT Publication No WO 91/19735), encoded peptides (PCT Publication WO 93/20242), random bio-oligomers (PCT Publication WO 92/00091), benzodiazepines (U.S. Pat. No. 5,288,514), diversomers such as hydantoins, benzodiazepines and dipeptides (Hobbs, et al. (1993) Proc. Nat. Acad. Sci. USA 90:6909-6913), vinylogous polypeptides (Hagihara, et al. (1992) J. Amer. Chem. Soc. 114:6568), nonpeptidal peptidomimetics with a Beta-D-Glucose scaffolding (Hirschmann, et

al. (1992) J. Amer. Chem. Soc. 114:9217-9218), analogous organic syntheses of small compound libraries (Chen, et al. (1994) J. Amer. Chem. Soc. 116:2661), oligocarbamates (Cho, et al. (1993) Science 261:1303), and/or peptidyl phosphonates (Campbell, et al. (1994) J. Org. Chem. 59:658). See, generally, Gordon, et al. (1994) J. Med. Chem. 37:1385, nucleic acid libraries (see, e.g., Stratagene, Corp.), peptide nucleic acid libraries (see, e.g., U.S. Patent 5,539,083), antibody libraries (see, e.g., Vaughn, et al. (1996) Nature Biotechnology 14(3):309-314, and PCT/US96/10287), carbohydrate libraries (see, e.g., Liang, et al. (1996) Science 274:1520-1522, and U.S. Patent No. 5,593,853), and small organic molecule libraries (see, e.g., benzodiazepines, Baum (1993) C&EN, Jan 18, page 33; isoprenoids, U.S. Patent No. 5,569,588; thiazolidinones and metathiazanones, U.S. Patent No. 5,549,974; pyrrolidines, U.S. Patent Nos. 5,525,735 and 5,519,134; morpholino compounds, U.S. Patent No. 5,506.337; benzodiazepines, U.S. Patent No. 5,288,514; and the like).

Devices for the preparation of combinatorial libraries are commercially available (see, e.g., 357 MPS, 390 MPS, Advanced Chem Tech, Louisville KY, Symphony, Rainin, Woburn, MA, 433A Applied Biosystems, Foster City, CA, 9050 Plus, Millipore, Bedford, MA).

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A number of well known robotic systems have also been developed for solution phase chemistries. These systems include automated workstations like the automated synthesis apparatus developed by Takeda Chemical Industries, LTD. (Osaka, Japan) and many robotic systems utilizing robotic arms (Zymate II, Zymark Corporation, Hopkinton, Mass.; Orca, Hewlett-Packard, Palo Alto, Calif.), which mimic the manual synthetic operations performed by a chemist. The above devices, with appropriate modification, are suitable for use with the present invention. In addition, numerous combinatorial libraries are themselves commercially available (see, e.g., ComGenex, Princeton, N.J., Asinex, Moscow, Ru, Tripos, Inc., St. Louis, MO, ChemStar, Ltd, Moscow, RU, 3D Pharmaceuticals, Exton, PA, Martek Biosciences, Columbia, MD, etc.).

The assays to identify modulators are amenable to high throughput screening.

Preferred assays thus detect modulation of lung cancer gene transcription, polypeptide expression, and polypeptide activity.

High throughput assays for evaluating the presence, absence, quantification, or other properties of particular nucleic acids or protein products are well known to those of skill in the art. Similarly, binding assays and reporter gene assays are similarly well known. Thus, e.g., U.S. Patent No. 5,559,410 discloses high throughput screening methods for proteins,

U.S. Patent No. 5,585,639 discloses high throughput screening methods for nucleic acid binding (i.e., in arrays), while U.S. Patent Nos. 5,576,220 and 5,541,061 disclose high throughput methods of screening for ligand/antibody binding.

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In addition, high throughput screening systems are commercially available (see, e.g., Zymark Corp., Hopkinton, MA; Air Technical Industries, Mentor, OH; Beckman Instruments, Inc. Fullerton, CA; Precision Systems, Inc., Natick, MA, etc.). These systems typically automate procedures, including sample and reagent pipetting, liquid dispensing, timed incubations, and final readings of the microplate in detector(s) appropriate for the assay. These configurable systems provide high throughput and rapid start up as well as a high degree of flexibility and customization. The manufacturers of such systems provide detailed protocols for various high throughput systems. Thus, e.g., Zymark Corp. provides technical bulletins describing screening systems for detecting the modulation of gene transcription, ligand binding, and the like.

In one embodiment, modulators are proteins, often naturally occurring proteins or fragments of naturally occurring proteins. Thus, e.g., cellular extracts containing proteins, or random or directed digests of proteinaceous cellular extracts, may be used. In this way libraries of proteins may be made for screening in the methods of the invention. Particularly preferred in this embodiment are libraries of bacterial, fungal, viral, and mammalian proteins, with the latter being preferred, and human proteins being especially preferred. Particularly useful test compound will be directed to the class of proteins to which the target belongs, e.g., substrates for enzymes or ligands and receptors.

In a preferred embodiment, modulators are peptides of from about 5 to about 30 amino acids, with from about 5 to about 20 amino acids being preferred, and from about 7 to about 15 being particularly preferred. The peptides may be digests of naturally occurring proteins, random peptides, or "biased" random peptides. By "randomized" or grammatical equivalents herein is meant that the nucleic acid or peptide consists of essentially random sequences of nucleotides and amino acids, respectively. Since these random peptides (or nucleic acids, discussed below) are often chemically synthesized, they may incorporate a nucleotide or amino acid at any position. The synthetic process can be designed to generate randomized proteins or nucleic acids, to allow the formation of all or most of the possible combinations over the length of the sequence, thus forming a library of randomized candidate bioactive proteinaceous agents.

In one embodiment, the library is fully randomized, with no sequence preferences or constants at any position. In a preferred embodiment, the library is biased. That is, some positions within the sequence are either held constant, or are selected from a limited number of possibilities. In a preferred embodiment, the nucleotides or amino acid residues are randomized within a defined class, e.g., of hydrophobic amino acids, hydrophilic residues, sterically biased (either small or large) residues, towards the creation of nucleic acid binding domains, the creation of cysteines, for cross-linking, prolines for SH-3 domains, serines, threonines, tyrosines or histidines for phosphorylation sites, etc.

Modulators of lung cancer can also be nucleic acids, as defined above.

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As described above generally for proteins, nucleic acid modulating agents may be naturally occurring nucleic acids, random nucleic acids, or "biased" random nucleic acids. Digests of procaryotic or eucaryotic genomes may be used as is outlined above for proteins.

In a preferred embodiment, the candidate compounds are organic chemical moieties, a wide variety of which are available in the literature.

After a candidate agent has been added and the cells allowed to incubate for some period of time, the sample containing a target sequence is analyzed. If required, the target sequence is prepared using known techniques. For example, the sample may be treated to lyse the cells, using known lysis buffers, electroporation, etc., with purification and/or amplification such as PCR performed as appropriate. For example, an *in vitro* transcription with labels covalently attached to the nucleotides is performed. Generally, the nucleic acids are labeled with biotin-FITC or PE, or with cy3 or cy5.

In a preferred embodiment, the target sequence is labeled with, e.g., a fluorescent, a chemiluminescent, a chemical, or a radioactive signal, to provide a means of detecting the target sequence's specific binding to a probe. The label also can be an enzyme, such as, alkaline phosphatase or horseradish peroxidase, which when provided with an appropriate substrate produces a product that can be detected. Alternatively, the label can be a labeled compound or small molecule, such as an enzyme inhibitor, that binds but is not catalyzed or altered by the enzyme. The label also can be a moiety or compound, such as, an epitope tag or biotin which specifically binds to streptavidin. For the example of biotin, the streptavidin is labeled as described above, thereby, providing a detectable signal for the bound target sequence. Unbound labeled streptavidin is typically removed prior to analysis.

Nucleic acid assays can be direct hybridization assays or can comprise "sandwich assays", which include the use of multiple probes, as is generally outlined in U.S. Patent Nos.

WO 02/086443

PCT/US02/12476
5,681,702, 5,597,909, 5,545,730, 5,594,117, 5,591,584, 5,571,670, 5,580,731, 5,571,670,
5,591,584, 5,624,802, 5,635,352, 5,594,118, 5,359,100, 5,124,246 and 5,681,697, all of which are hereby incorporated by reference. In this embodiment, in general, the target nucleic acid is prepared as outlined above, and then added to the biochip comprising a plurality of nucleic acid probes, under conditions that allow the formation of a hybridization complex.

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A variety of hybridization conditions may be used in the present invention, including high, moderate and low stringency conditions as outlined above. The assays are generally run under stringency conditions which allow formation of the label probe hybridization complex only in the presence of target. Stringency can be controlled by altering a step parameter that is a thermodynamic variable, including, but not limited to, temperature, formamide concentration, salt concentration, chaotropic salt concentration, pH, organic solvent concentration, etc.

These parameters may also be used to control non-specific binding, as is generally outlined in U.S. Patent No. 5,681,697. Thus it may be desirable to perform certain steps at higher stringency conditions to reduce non-specific binding.

The reactions outlined herein may be accomplished in a variety of ways. Components of the reaction may be added simultaneously, or sequentially, in different orders, with preferred embodiments outlined below. In addition, the reaction may include a variety of other reagents. These include salts, buffers, neutral proteins, e.g., albumin, detergents, etc. which may be used to facilitate optimal hybridization and detection, and/or reduce non-specific or background interactions. Reagents that otherwise improve the efficiency of the assay, such as protease inhibitors, nuclease inhibitors, anti-microbial agents, etc., may also be used as appropriate, depending on the sample preparation methods and purity of the target.

The assay data are analyzed to determine the expression levels, and changes in expression levels as between states, of individual genes, forming a gene expression profile.

Screens are performed to identify modulators of the lung cancer phenotype. In one embodiment, screening is performed to identify modulators that can induce or suppress a particular expression profile, thus preferably generating the associated phenotype. In another embodiment, e.g., for diagnostic applications, having identified differentially expressed genes important in a particular state, screens can be performed to identify modulators that alter expression of individual genes. In an another embodiment, screening is performed to identify modulators that alter a biological function of the expression product of a differentially expressed gene. Again, having identified the importance of a gene in a particular state,

screens are performed to identify agents that bind and/or modulate the biological activity of the gene product, or evaluate genetic polymorphisms.

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Genes can be screened for those that are induced in response to a candidate agent. After identifying a modulator based upon its ability to suppress a lung cancer expression pattern leading to a normal expression pattern, or to modulate a single lung cancer gene expression profile so as to mimic the expression of the gene from normal tissue, a screen as described above can be performed to identify genes that are specifically modulated in response to the agent. Comparing expression profiles between normal tissue and agent treated lung cancer tissue reveals genes that are not expressed in normal tissue or lung cancer tissue, but are expressed in agent treated tissue. These agent-specific sequences can be identified and used by methods described herein for lung cancer genes or proteins. In particular these sequences and the proteins they encode find use in marking or identifying agent treated cells. In addition, antibodies can be raised against the agent induced proteins and used to target novel therapeutics to the treated lung cancer tissue sample.

Thus, in one embodiment, a test compound is administered to a population of lung cancer cells, that have an associated lung cancer expression profile. By "administration" or "contacting" herein is meant that the candidate agent is added to the cells in such a manner as to allow the agent to act upon the cell, whether by uptake and intracellular action, or by action at the cell surface. In some embodiments, nucleic acid encoding a proteinaceous candidate agent (i.e., a peptide) may be put into a viral construct such as an adenoviral or retroviral construct, and added to the cell, such that expression of the peptide agent is accomplished, e.g., PCT US97/01019. Regulatable gene therapy systems can also be used.

Once a test compound has been administered to the cells, the cells can be washed if desired and are allowed to incubate under preferably physiological conditions for some period of time. The cells are then harvested and a new gene expression profile is generated, as outlined herein.

Thus, e.g., lung cancer or non-malignant tissue may be screened for agents that modulate, e.g., induce or suppress a lung cancer phenotype. A change in at least one gene, preferably many, of the expression profile indicates that the agent has an effect on lung cancer activity. By defining such a signature for the lung cancer phenotype, screens for new drugs that alter the phenotype can be devised. With this approach, the drug target need not be known and need not be represented in the original expression screening platform, nor does the level of transcript for the target protein need to change.

Measure of lung cancer polypeptide activity, or of lung cancer or the lung cancer phenotype can be performed using a variety of assays. For example, the effects of the test compounds upon the function of the metastatic polypeptides can be measured by examining parameters described above. A suitable physiological change that affects activity can be used to assess the influence of a test compound on the polypeptides of this invention. When the functional consequences are determined using intact cells or animals, one can also measure a variety of effects such as, in the case of lung cancer associated with tumors, tumor growth, tumor metastasis, neovascularization, hormone release, transcriptional changes to both known and uncharacterized genetic markers (e.g., northern blots), changes in cell metabolism such as cell growth or pH changes, and changes in intracellular second messengers such as cGMP. In the assays of the invention, mammalian lung cancer polypeptide is typically used, e.g., mouse, preferably human.

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Assays to identify compounds with modulating activity can be performed *in vitro*. For example, a lung cancer polypeptide is first contacted with a potential modulator and incubated for a suitable amount of time, e.g., from 0.5 to 48 hours. In one embodiment, the lung cancer polypeptide levels are determined *in vitro* by measuring the level of protein or mRNA. The level of protein is typically measured using immunoassays such as western blotting, ELISA and the like with an antibody that selectively binds to the lung cancer polypeptide or a fragment thereof. For measurement of mRNA, amplification, e.g., using PCR, LCR, or hybridization assays, e.g., northern hybridization, RNAse protection, dot blotting, are preferred. The level of protein or mRNA is typically detected using directly or indirectly labeled detection agents, e.g., fluorescently or radioactively labeled nucleic acids, radioactively or enzymatically labeled antibodies, and the like, as described herein.

Alternatively, a reporter gene system can be devised using a lung cancer protein promoter operably linked to a reporter gene such as luciferase, green fluorescent protein, CAT, or β -gal. The reporter construct is typically transfected into a cell. After treatment with a potential modulator, the amount of reporter gene transcription, translation, or activity is measured according to standard techniques known to those of skill in the art.

In a preferred embodiment, as outlined above, screens may be done on individual genes and gene products (proteins). That is, having identified a particular differentially expressed gene as important in a particular state, screening of modulators of the expression of the gene product itself can be done. The gene products of differentially expressed

genes are sometimes referred to herein as "lung cancer proteins." The lung cancer protein may be a fragment, or alternatively, be the full length protein to a fragment shown herein.

In one embodiment, screening for modulators of expression of specific genes is performed. Typically, the expression of only one or a few genes are evaluated. In another embodiment, screens are designed to first find compounds that bind to differentially expressed proteins. These compounds are then evaluated for the ability to modulate differentially expressed activity. Moreover, once initial candidate compounds are identified, variants can be further screened to better evaluate structure activity relationships.

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In a preferred embodiment, binding assays are done. In general, purified or isolated gene product is used; that is, the gene products of one or more differentially expressed nucleic acids are made. For example, antibodies are generated to the protein gene products, and standard immunoassays are run to determine the amount of protein present. Alternatively, cells comprising the lung cancer proteins can be used in the assays.

Thus, in a preferred embodiment, the methods comprise combining a lung cancer protein and a candidate compound, and determining the binding of the compound to the lung cancer protein. Preferred embodiments utilize the human lung cancer protein, although other mammalian proteins may also be used, e.g., for the development of animal models of human disease. In some embodiments, as outlined herein, variant or derivative lung cancer proteins may be used.

Generally, in a preferred embodiment of the methods herein, the lung cancer protein or the candidate agent is non-diffusably bound to an insoluble support, preferably having isolated sample receiving areas (e.g., a microtiter plate, an array, etc.). The insoluble supports may be made of a composition to which the compositions can be bound, is readily separated from soluble material, and is otherwise compatible with the overall method of screening. The surface of such supports may be solid or porous and of a convenient shape. Examples of suitable insoluble supports include microtiter plates, arrays, membranes and beads. These are typically made of glass, plastic (e.g., polystyrene), polysaccharides, nylon or nitrocellulose, teflonTM, etc. Microtiter plates and arrays are especially convenient because a large number of assays can be carried out simultaneously, using small amounts of reagents and samples. The particular manner of binding of the composition is typically not crucial so long as it is compatible with the reagents and overall methods of the invention, maintains the activity of the composition, and is nondiffusable. Preferred methods of binding include the use of antibodies (which do not sterically block either the ligand binding site or activation

sequence when the protein is bound to the support), direct binding to "sticky" or ionic supports, chemical crosslinking, the synthesis of the protein or agent on the surface, etc. Following binding of the protein or agent, excess unbound material is removed by washing. The sample receiving areas may then be blocked through incubation with bovine serum albumin (BSA), casein or other innocuous protein or other moiety.

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In a preferred embodiment, the lung cancer protein is bound to the support, and a test compound is added to the assay. Alternatively, the candidate agent is bound to the support and the lung cancer protein is added. Novel binding agents include specific antibodies, nonnatural binding agents identified in screens of chemical libraries, peptide analogs, etc. Of particular interest are screening assays for agents that have a low toxicity for human cells. A wide variety of assays may be used for this purpose, including labeled in vitro protein-protein binding assays, electrophoretic mobility shift assays, immunoassays for protein binding, functional assays (phosphorylation assays, etc.) and the like.

The determination of the binding of the test modulating compound to the lung cancer protein may be done in a number of ways. In a preferred embodiment, the compound is labeled, and binding determined directly, e.g., by attaching all or a portion of the lung cancer protein to a solid support, adding a labeled candidate agent (e.g., a fluorescent label), washing off excess reagent, and determining whether the label is present on the solid support. Various blocking and washing steps may be utilized as appropriate.

In some embodiments, only one of the components is labeled, e.g., the proteins (or proteinaceous candidate compounds) can be labeled. Alternatively, more than one component can be labeled with different labels, e.g., 125I for the proteins and a fluorophor for the compound. Proximity reagents, e.g., quenching or energy transfer reagents are also useful.

In one embodiment, the binding of the test compound is determined by competitive binding assay. The competitor may be a binding moiety known to bind to the target molecule (i.e., a lung cancer protein), such as an antibody, peptide, binding partner, ligand, etc. Under certain circumstances, there may be competitive binding between the compound and the binding moiety, with the binding moiety displacing the compound. In one embodiment, the 30 test compound is labeled. Either the compound, or the competitor, or both, is added first to the protein for a time sufficient to allow binding, if present. Incubations may be performed at a temperature which facilitates optimal activity, typically between 4 and 40° C. Incubation periods are typically optimized, e.g., to facilitate rapid high throughput screening. Typically

between 0.1 and 1 hour will be sufficient. Excess reagent is generally removed or washed away. The second component is then added, and the presence or absence of the labeled component is followed, to indicate binding.

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In a preferred embodiment, the competitor is added first, followed by a test compound. Displacement of the competitor is an indication that the test compound is binding to the lung cancer protein and thus is capable of binding to, and potentially modulating, the activity of the lung cancer protein. In this embodiment, either component can be labeled. Thus, e.g., if the competitor is labeled, the presence of label in the wash solution indicates displacement by the agent. Alternatively, if the test compound is labeled, the presence of the label on the support indicates displacement.

In an alternative embodiment, the test compound is added first, with incubation and washing, followed by the competitor. The absence of binding by the competitor may indicate that the test compound is bound to the lung cancer protein with a higher affinity. Thus, if the test compound is labeled, the presence of the label on the support, coupled with a lack of competitor binding, may indicate that the test compound is capable of binding to the lung cancer protein.

In a preferred embodiment, the methods comprise differential screening to identity agents that are capable of modulating the activity of the lung cancer proteins. In one embodiment, the methods comprise combining a lung cancer protein and a competitor in a first sample. A second sample comprises a test compound, a lung cancer protein, and a competitor. The binding of the competitor is determined for both samples, and a change, or difference in binding between the two samples indicates the presence of an agent capable of binding to the lung cancer protein and potentially modulating its activity. That is, if the binding of the competitor is different in the second sample relative to the first sample, the agent is capable of binding to the lung cancer protein.

Alternatively, differential screening is used to identify drug candidates that bind to the native lung cancer protein, but cannot bind to modified lung cancer proteins. The structure of the lung cancer protein may be modeled, and used in rational drug design to synthesize agents that interact with that site. Drug candidates that affect the activity of a lung cancer protein are also identified by screening drugs for the ability to either enhance or reduce the activity of the protein.

Positive controls and negative controls may be used in the assays. Preferably control and test samples are performed in at least triplicate to obtain statistically significant results.

Incubation of all samples is for a time sufficient for the binding of the agent to the protein. Following incubation, samples are washed free of non-specifically bound material and the amount of bound, generally labeled agent determined. For example, where a radiolabel is employed, the samples may be counted in a scintillation counter to determine the amount of bound compound.

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A variety of other reagents may be included in the screening assays. These include reagents like salts, neutral proteins, e.g., albumin, detergents, etc. which may be used to facilitate optimal protein-protein binding and/or reduce non-specific or background interactions. Also reagents that otherwise improve the efficiency of the assay, such as protease inhibitors, nuclease inhibitors, anti-microbial agents, etc., may be used. The mixture of components may be added in an order that provides for the requisite binding.

In a preferred embodiment, the invention provides methods for screening for a compound capable of modulating the activity of a lung cancer protein. The methods comprise adding a test compound, as defined above, to a cell comprising lung cancer proteins. Preferred cell types include almost any cell. The cells contain a recombinant nucleic acid that encodes a lung cancer protein. In a preferred embodiment, a library of candidate agents are tested on a plurality of cells.

In one aspect, the assays are evaluated in the presence or absence or previous or subsequent exposure of physiological signals, e.g., hormones, antibodies, peptides, antigens, cytokines, growth factors, action potentials, pharmacological agents including chemotherapeutics, radiation, carcinogenics, or other cells (e.g., cell-cell contacts). In another example, the determinations are determined at different stages of the cell cycle process.

In this way, compounds that modulate lung cancer agents are identified. Compounds with pharmacological activity are able to enhance or interfere with the activity of the lung cancer protein. Once identified, similar structures are evaluated to identify critical structural feature of the compound.

In one embodiment, a method of inhibiting lung cancer cell division is provided. The method comprises administration of a lung cancer inhibitor. In another embodiment, a method of inhibiting lung cancer is provided. The method may comprise administration of a lung cancer inhibitor. In a further embodiment, methods of treating cells or individuals with lung cancer are provided, e.g., comprising administration of a lung cancer inhibitor.

In one embodiment, a lung cancer inhibitor is an antibody as discussed above. In another embodiment, the lung cancer inhibitor is an antisense molecule.

A variety of cell growth, proliferation, viability, and metastasis assays are known to those of skill in the art, as described below.

Soft agar growth or colony formation in suspension

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Normal cells require a solid substrate to attach and grow. When the cells are transformed, they lose this phenotype and grow detached from the substrate. For example, transformed cells can grow in stirred suspension culture or suspended in semi-solid media, such as semi-solid or soft agar. The transformed cells, when transfected with tumor suppressor genes, regenerate normal phenotype and require a solid substrate to attach and grow. Soft agar growth or colony formation in suspension assays can be used to identify modulators of lung cancer sequences, which when expressed in host cells, inhibit abnormal cellular proliferation and transformation. A therapeutic compound would reduce or eliminate the host cells' ability to grow in stirred suspension culture or suspended in semi-solid media, such as semi-solid or soft.

Techniques for soft agar growth or colony formation in suspension assays are described in Freshney (1994) <u>Culture of Animal Cells a Manual of Basic Technique</u> (3rd ed.), herein incorporated by reference. See also, the methods section of Garkavtsev, et al. (1996), *supra*, herein incorporated by reference.

Contact inhibition and density limitation of growth

Normal cells typically grow in a flat and organized pattern in a petri dish until they touch other cells. When the cells touch one another, they are contact inhibited and stop growing. When cells are transformed, however, the cells are not contact inhibited and continue to grow to high densities in disorganized foci. Thus, the transformed cells grow to a higher saturation density than normal cells. This can be detected morphologically by the formation of a disoriented monolayer of cells or rounded cells in foci within the regular pattern of normal surrounding cells. Alternatively, labeling index with (³H)-thymidine at saturation density can be used to measure density limitation of growth. See Freshney (1994), supra. The transformed cells, when transfected with tumor suppressor genes, regenerate a normal phenotype and become contact inhibited and would grow to a lower density.

In this assay, labeling index with (³H)-thymidine at saturation density is a preferred method of measuring density limitation of growth. Transformed host cells are transfected with a lung cancer-associated sequence and are grown for 24 hours at saturation density in

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non-limiting medium conditions. The percentage of cells labeling with (³H)-thymidine is determined autoradiographically. See, Freshney (1994), *supra*.

Growth factor or serum dependence

Transformed cells typically have a lower serum dependence than their normal counterparts (see, e.g., Temin (1966) <u>J. Natl. Cancer Insti.</u> 37:167-175; Eagle, et al. (1970) <u>J. Exp. Med.</u> 131:836-879); Freshney, *supra*. This is in part due to release of various growth factors by the transformed cells. Growth factor or serum dependence of transformed host cells can be compared with that of control.

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Tumor specific markers levels

Tumor cells release an increased amount of certain factors (hereinafter "tumor specific markers") than their normal counterparts. For example, plasminogen activator (PA) is released from human glioma at a higher level than from normal brain cells (see, e.g., Gullino, "Angiogenesis, tumor vascularization, and potential interference with tumor growth" in Mihich (ed. 1985) <u>Biological Responses in Cancer</u>, pp. 178-184). Similarly, Tumor angiogenesis factor (TAF) is released at a higher level in tumor cells than their normal counterparts. See, e.g., Folkman (1992) "Angiogenesis and Cancer" in <u>Sem Cancer Biol.</u>).

Various techniques which measure the release of these factors are described in Freshney (1994), *supra*. Also, see, Unkeless, et al. (1974) <u>J. Biol. Chem.</u> 249:4295-4305; Strickland and Beers (1976) <u>J. Biol. Chem.</u> 251:5694-5702; Whur, et al. (1980) <u>Br. J. Cancer</u> 42:305-312; Gullino, "Angiogenesis, tumor vascularization, and potential interference with tumor growth" in Mihich (ed. 1985) <u>Biological Responses in Cancer</u>, pp. 178-184; Freshney <u>Anticancer Res.</u> 5:111-130 (1985).

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Invasiveness into Matrigel

The degree of invasiveness into Matrigel or some other extracellular matrix constituent can be used as an assay to identify compounds that modulate lung cancer-associated sequences. Tumor cells exhibit a good correlation between malignancy and invasiveness of cells into Matrigel or some other extracellular matrix constituent. In this assay, tumorigenic cells are typically used as host cells. Expression of a tumor suppressor gene in these host cells would decrease invasiveness of the host cells.

Techniques described in Freshney (1994), *supra*, can be used. Briefly, the level of invasion of host cells can be measured by using filters coated with Matrigel or some other extracellular matrix constituent. Penetration into the gel, or through to the distal side of the filter, is rated as invasiveness, and rated histologically by number of cells and distance moved, or by prelabeling the cells with ¹²⁵I and counting the radioactivity on the distal side of the filter or bottom of the dish. See, e.g., Freshney (1984), *supra*.

Tumor growth in vivo

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Effects of lung cancer-associated sequences on cell growth can be tested in transgenic or immune-suppressed mice. Knock-out transgenic mice can be made, in which the lung cancer gene is disrupted or in which a lung cancer gene is inserted. Knock-out transgenic mice can be made by insertion of a marker gene or other heterologous gene into the endogenous lung cancer gene site in the mouse genome via homologous recombination. Such mice can also be made by substituting the endogenous lung cancer gene with a mutated version of the lung cancer gene, or by mutating the endogenous lung cancer gene, e.g., by exposure to carcinogens.

A DNA construct is introduced into the nuclei of embryonic stem cells. Cells containing the newly engineered genetic lesion are injected into a host mouse embryo, which is re-implanted into a recipient female. Some of these embryos develop into chimeric mice that possess germ cells partially derived from the mutant cell line. Therefore, by breeding the chimeric mice it is possible to obtain a new line of mice containing the introduced genetic lesion (see, e.g., Capecchi, et al. (1989) Science 244:1288). Chimeric targeted mice can be derived according to Hogan, et al. (1988) Manipulating the Mouse Embryo: A Laboratory Manual, Cold Spring Harbor Laboratory and Robertson (ed. 1987) Teratocarcinomas and Embryonic Stem Cells: A Practical Approach, , IRL Press, Washington, D.C.

Alternatively, various immune-suppressed or immune-deficient host animals can be used. For example, genetically athymic "nude" mouse (see, e.g., Giovanella, et al. (1974) <u>J. Natl. Cancer Inst.</u> 52:921), a SCID mouse, a thymectomized mouse, or an irradiated mouse (see, e.g., Bradley, et al. (1978) <u>Br. J. Cancer</u> 38:263; Selby, et al. (1980) <u>Br. J. Cancer</u> 41:52) can be used as a host. Transplantable tumor cells (typically about 10⁶ cells) injected into isogenic hosts will produce invasive tumors in a high proportions of cases, while normal cells of similar origin will not. In hosts which developed invasive tumors, cells expressing a lung cancer-associated sequences are injected subcutaneously. After a suitable length of time,

preferably 4-8 weeks, tumor growth is measured (e.g., by volume or by its two largest dimensions) and compared to the control. Tumors that have statistically significant reduction (using, e.g., Student's T test) are said to have inhibited growth.

5 Polynucleotide modulators of lung cancer

Antisense and RNAi Polynucleotides

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In certain embodiments, the activity of a lung cancer-associated protein is downregulated, or entirely inhibited, by the use of antisense or an inhibitory polynucleotide, i.e., a nucleic acid complementary to, and which can preferably hybridize specifically to, a coding mRNA nucleic acid sequence, e.g., a lung cancer protein mRNA, or a subsequence thereof. Binding of the antisense polynucleotide to the mRNA reduces the translation and/or stability of the mRNA.

In the context of this invention, antisense polynucleotides can comprise naturally-occurring nucleotides, or synthetic species formed from naturally-occurring subunits or their close homologs. Antisense polynucleotides may also have altered sugar moieties or intersugar linkages. Exemplary among these are the phosphorothioate and other sulfur containing species which are known for use in the art. Analogs are comprehended by this invention so long as they function effectively to hybridize with the lung cancer protein mRNA. See, e.g., Isis Pharmaceuticals, Carlsbad, CA; Sequitor, Inc., Natick, MA.

Such antisense polynucleotides can readily be synthesized using recombinant means, or can be synthesized *in vitro*. Equipment for such synthesis is sold by several vendors, including Applied Biosystems. The preparation of other oligonucleotides such as phosphorothioates and alkylated derivatives is also well known to those of skill in the art.

Antisense molecules as used herein include antisense or sense oligonucleotides.

Sense oligonucleotides can, e.g., be employed to block transcription by binding to the antisense strand. The antisense and sense oligonucleotide comprise a single-stranded nucleic acid sequence (either RNA or DNA) capable of binding to target mRNA (sense) or DNA (antisense) sequences for lung cancer molecules. A preferred antisense molecule is for a lung cancer sequence in the tables, or for a ligand or activator thereof. Antisense or sense oligonucleotides, according to the present invention, comprise a fragment generally at least about 14 nucleotides, preferably from about 14 to 30 nucleotides. The ability to derive an antisense or a sense oligonucleotide, based upon a cDNA sequence encoding a given protein

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is described in, e.g., Stein and Cohen (1988) Cancer Res. 48:2659 and van der Krol, et al.

(1988) BioTechniques 6:958).

RNA interference is a mechanism to suppress gene expression in a sequence specific manner. See, e.g., Brumelkamp, et al. (2002) Sciencexpress (21March2002); Sharp (1999) Genes Dev. 13:139-141; and Cathew (2001) Curr. Op. Cell Biol. 13:244-248. In mammalian cells, short, e.g., 21 nt, double stranded small interfering RNAs (siRNA) have been shown to be effective at inducing an RNAi response. See, e.g., Elbashir, et al. (2001) Nature 411:494-498. The mechanism may be used to downregulate expression levels of identified genes, e.g., treatment of or validation of relevance to disease.

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Ribozymes

In addition to antisense polynucleotides, ribozymes can be used to target and inhibit transcription of lung cancer-associated nucleotide sequences. A ribozyme is an RNA molecule that catalytically cleaves other RNA molecules. Different kinds of ribozymes have been described, including group I ribozymes, hammerhead ribozymes, hairpin ribozymes, RNase P, and axhead ribozymes (see, e.g., Castanotto, et al. (1994) Adv. in Pharmacology 25: 289-317 for a general review of the properties of different ribozymes).

The general features of hairpin ribozymes are described, e.g., in Hampel, et al. (1990) Nucl. Acids Res. 18:299-304; European Patent Publication No. 0 360 257; U.S. Patent No. 5,254,678. Methods of preparing are well known to those of skill in the art (see, e.g., WO 94/26877; Ojwang, et al. (1993) Proc. Natl. Acad. Sci. USA 90:6340-6344; Yamada, et al. (1994) Human Gene Therapy 1:39-45; Leavitt, et al. (1995) Proc. Natl. Acad. Sci. USA 92:699-703; Leavitt, et al. (1994) Human Gene Therapy 5:1151-120; and Yamada, et al. (1994) Virology 205: 121-126).

Polynucleotide modulators of lung cancer may be introduced into a cell containing the

target nucleotide sequence by formation of a conjugate with a ligand binding molecule, as described in WO 91/04753. Suitable ligand binding molecules include, but are not limited to, cell surface receptors, growth factors, other cytokines, or other ligands that bind to cell surface receptors. Preferably, conjugation of the ligand binding molecule does not substantially interfere with the ability of the ligand binding molecule to bind to its corresponding molecule or receptor, or block entry of the sense or antisense oligonucleotide or its conjugated version into the cell. Alternatively, a polynucleotide modulator of lung cancer may be introduced into a cell containing the target nucleic acid sequence, e.g., by

formation of an polynucleotide-lipid complex, as described in WO 90/10448. It is understood that the use of antisense molecules or knock out and knock in models may also be used in screening assays as discussed above, in addition to methods of treatment.

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Thus, in one embodiment, methods of modulating lung cancer in cells or organisms are provided. In one embodiment, the methods comprise administering to a cell an anti-lung cancer antibody that reduces or eliminates the biological activity of an endogenous lung cancer protein. Alternatively, the methods comprise administering to a cell or organism a recombinant nucleic acid encoding a lung cancer protein. This may be accomplished in any number of ways. In a preferred embodiment, e.g., when the lung cancer sequence is down-regulated in lung cancer, such state may be reversed by increasing the amount of lung cancer gene product in the cell. This can be accomplished, e.g., by overexpressing the endogenous lung cancer gene or administering a gene encoding the lung cancer sequence, using known gene-therapy techniques. In a preferred embodiment, the gene therapy techniques include the incorporation of the exogenous gene using enhanced homologous recombination (EHR), e.g., as described in PCT/US93/03868, hereby incorporated by reference in its entirety. Alternatively, e.g., when the lung cancer sequence is up-regulated in lung cancer, the activity of the endogenous lung cancer gene is decreased, e.g., by the administration of a lung cancer antisense or RNAi nucleic acid.

In one embodiment, the lung cancer proteins of the present invention may be used to generate polyclonal and monoclonal antibodies to lung cancer proteins. Similarly, the lung cancer proteins can be coupled, using standard technology, to affinity chromatography columns. These columns may then be used to purify lung cancer antibodies useful for production, diagnostic, or therapeutic purposes. In a preferred embodiment, the antibodies are generated to epitopes unique to a lung cancer protein; that is, the antibodies show little or no cross-reactivity to other proteins. The lung cancer antibodies may be coupled to standard affinity chromatography columns and used to purify lung cancer proteins. The antibodies may also be used as blocking polypeptides, as outlined above, since they will specifically bind to the lung cancer protein.

30 Methods of identifying variant lung cancer-associated sequences

Without being bound by theory, expression of various lung cancer sequences is correlated with lung cancer. Accordingly, disorders based on mutant or variant lung cancer genes may be determined. In one embodiment, the invention provides methods for

identifying cells containing variant lung cancer genes, e.g., determining all or part of the sequence of at least one endogenous lung cancer genes in a cell. In a preferred embodiment, the invention provides methods of identifying the lung cancer genotype of an individual, e.g., determining all or part of the sequence of at least one lung cancer gene of the individual. This is generally done in at least one tissue of the individual, and may include the evaluation of a number of tissues or different samples of the same tissue. The method may include comparing the sequence of the sequenced lung cancer gene to a known lung cancer gene, i.e., a wild-type gene.

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The sequence of all or part of the lung cancer gene can then be compared to the sequence of a known lung cancer gene to determine if any differences exist. This can be done using known homology programs, such as Bestfit, etc. In a preferred embodiment, the presence of a difference in the sequence between the lung cancer gene of the patient and the known lung cancer gene correlates with a disease state or a propensity for a disease state, as outlined herein.

In a preferred embodiment, the lung cancer genes are used as probes to determine the number of copies of the lung cancer gene in the genome.

In another preferred embodiment, the lung cancer genes are used as probes to determine the chromosomal localization of the lung cancer genes. Information such as chromosomal localization finds use in providing a diagnosis or prognosis in particular when chromosomal abnormalities such as translocations, and the like are identified in the lung cancer gene locus.

Administration of pharmaceutical and vaccine compositions

In one embodiment, a therapeutically effective dose of a lung cancer protein or modulator thereof, is administered to a patient. By "therapeutically effective dose" herein is meant a dose that produces effects for which it is administered. The exact dose will depend on the purpose of the treatment, and will be ascertainable by one skilled in the art using known techniques (e.g., Ansel, et al. (1992) Pharmaceutical Dosage Forms and Drug Delivery; Lieberman, Pharmaceutical Dosage Forms (vols. 1-3), Dekker, ISBN 0824770846, 082476918X, 0824712692, 0824716981; Lloyd (1999) The Art, Science and Technology of Pharmaceutical Compounding; and Pickar (1999) Dosage Calculations). Adjustments for lung cancer degradation, systemic versus localized delivery, and rate of new protease synthesis, as well as the age, body weight, general health, sex, diet, time of administration,

drug interaction and the severity of the condition may be necessary, and will be ascertainable with routine experimentation by those skilled in the art.

A "patient" for the purposes of the present invention includes both humans and other animals, particularly mammals. Thus the methods are applicable to both human therapy and veterinary applications. In the preferred embodiment the patient is a mammal, preferably a primate, and in the most preferred embodiment the patient is human.

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The administration of the lung cancer proteins and modulators thereof of the present invention can be done in a variety of ways, including, but not limited to, orally, subcutaneously, intravenously, intranasally, transdermally, intraperitoneally, intramuscularly, intrapulmonary, vaginally, rectally, or intraocularly. In some instances, e.g., in the treatment of wounds and inflammation, the lung cancer proteins and modulators may be directly applied as a solution or spray.

The pharmaceutical compositions of the present invention comprise a lung cancer protein in a form suitable for administration to a patient. In the preferred embodiment, the pharmaceutical compositions are in a water soluble form, such as being present as pharmaceutically acceptable salts, which is meant to include both acid and base addition salts. "Pharmaceutically acceptable acid addition salt" refers to those salts that retain the biological effectiveness of the free bases and that are not biologically or otherwise undesirable, formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid and the like, and organic acids such as acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, p-toluenesulfonic acid, salicylic acid and the like. "Pharmaceutically acceptable base addition salts" include those derived from inorganic bases such as sodium, potassium, lithium, ammonium, calcium, magnesium, iron, zinc, copper, manganese, aluminum salts and the like. Particularly preferred are the ammonium, potassium, sodium, calcium, and magnesium salts. Salts derived from pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, and ethanolamine.

The pharmaceutical compositions may also include one or more of the following: carrier proteins such as serum albumin; buffers; fillers such as microcrystalline cellulose,

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coloring agents; and polyethylene glycol.

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The pharmaceutical compositions can be administered in a variety of unit dosage forms depending upon the method of administration. For example, unit dosage forms suitable for oral administration include, but are not limited to, powder, tablets, pills, capsules and lozenges. It is recognized that lung cancer protein modulators (e.g., antibodies, antisense constructs, ribozymes, small organic molecules, etc.) when administered orally, should be protected from digestion. This is typically accomplished either by complexing the molecule(s) with a composition to render it resistant to acidic and enzymatic hydrolysis, or by packaging the molecule(s) in an appropriately resistant carrier, such as a liposome or a protection barrier. Means of protecting agents from digestion are well known in the art.

The compositions for administration will commonly comprise a lung cancer protein modulator dissolved in a pharmaceutically acceptable carrier, preferably an aqueous carrier. A variety of aqueous carriers can be used, e.g., buffered saline and the like. These solutions are sterile and generally free of undesirable matter. These compositions may be sterilized by conventional, well known sterilization techniques. The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions such as pH adjusting and buffering agents, toxicity adjusting agents and the like, e.g., sodium acetate, sodium chloride, potassium chloride, calcium chloride, sodium lactate and the like. The concentration of active agent in these formulations can vary widely, and will be selected primarily based on fluid volumes, viscosities, body weight and the like in accordance with the particular mode of administration selected and the patient's needs (e.g., Remington's Pharmaceutical Science (15th ed., 1980) and Hardman, et al. (eds. 1996)

Goodman and Gilman: The Pharmacologial Basis of Therapeutics).

Thus, a typical pharmaceutical composition for intravenous administration would be about 0.1 to 10 mg per patient per day. Dosages from 0.1 up to about 100 mg per patient per day may be used, particularly when the drug is administered to a secluded site and not into the blood stream, such as into a body cavity or into a lumen of an organ. Substantially higher dosages are possible in topical administration. Actual methods for preparing parenterally administrable compositions will be known or apparent to those skilled in the art, e.g., Remington's Pharmaceutical Science and Goodman and Gilman, The Pharmacologial Basis of Therapeutics, supra.

The compositions containing modulators of lung cancer proteins can be administered for therapeutic or prophylactic treatments. In therapeutic applications, compositions are administered to a patient suffering from a disease (e.g., a cancer) in an amount sufficient to cure or at least partially arrest the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the disease and the general state of the patient's health. Single or multiple administrations of the compositions may be administered depending on the dosage and frequency as required and tolerated by the patient. In any event, the composition should provide a sufficient quantity of the agents of this invention to effectively treat the patient. An amount of modulator that is capable of preventing or slowing the development of cancer in a mammal is referred to as a "prophylactically effective dose." The particular dose required for a prophylactic treatment will depend upon the medical condition and history of the mammal, the particular cancer being prevented, as well as other factors such as age, weight, gender, administration route, efficiency, etc. Such prophylactic treatments may be used, e.g., in a mammal who has previously had cancer to prevent a recurrence of the cancer, or in a mammal who is suspected of having a significant likelihood of developing cancer based, at least in part, upon gene expression profiles. Vaccine strategies may be used, in either a DNA vaccine form, or protein vaccine.

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It will be appreciated that the present lung cancer protein-modulating compounds can be administered alone or in combination with additional lung cancer modulating compounds or with other therapeutic agent, e.g., other anti-cancer agents or treatments.

In numerous embodiments, one or more nucleic acids, e.g., polynucleotides comprising nucleic acid sequences set forth in the tables, such as antisense or RNAi polynucleotides or ribozymes, will be introduced into cells, *in vitro* or *in vivo*. The present invention provides methods, reagents, vectors, and cells useful for expression of lung cancer-associated polypeptides and nucleic acids using *in vitro* (cell-free), *ex vivo*, or *in vivo* (cell or organism-based) recombinant expression systems.

The particular procedure used to introduce the nucleic acids into a host cell for expression of a protein or nucleic acid is application specific. Many procedures for introducing foreign nucleotide sequences into host cells may be used. These include the use of calcium phosphate transfection, spheroplasts, electroporation, liposomes, microinjection, plasma vectors, viral vectors and other well known methods for introducing cloned genomic DNA, cDNA, synthetic DNA or other foreign genetic material into a host cell (see, e.g.,

Berger and Kimmel, <u>Guide to Molecular Cloning Techniques</u>, <u>Methods in Enzymology</u> volume 152 (Berger), Ausubel, et al. (eds. 1999) <u>Current Protocols</u> (supplemented through 1999), and Sambrook, et al. (1989) <u>Molecular Cloning - A Laboratory Manual</u> (2nd ed., Vol. 1-3).

In a preferred embodiment, lung cancer proteins and modulators are administered as therapeutic agents, and can be formulated as outlined above. Similarly, lung cancer genes (including both the full-length sequence, partial sequences, or regulatory sequences of the lung cancer coding regions) can be administered in a gene therapy application. These lung cancer genes can include antisense or inhibitory applications, e.g., as inhibitory RNA or gene therapy (e.g., for incorporation into the genome) or as antisense compositions.

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Lung cancer polypeptides and polynucleotides can also be administered as vaccine compositions to stimulate HTL, CTL, and antibody responses.. Such vaccine compositions can include, e.g., lipidated peptides (see, e.g., Vitiello, et al. (1995) J. Clin. Invest. 95:341), peptide compositions encapsulated in poly(DL-lactide-co-glycolide) ("PLG") microspheres (see, e.g., Eldridge, et al. (1991) Molec. Immunol. 28:287-294; Alonso, et al. (1994) Vaccine 12:299-306; Jones, et al. (1995) Vaccine 13:675-681), peptide compositions contained in immune stimulating complexes (ISCOMS) (see, e.g., Takahashi, et al. (1990) Nature 344:873-875; Hu, et al. (1998) Clin Exp Immunol. 113:235-243), multiple antigen peptide systems (MAPs) (see, e.g., Tam (1988) Proc. Natl. Acad. Sci. U.S.A. 85:5409-5413; Tam (1996) J. Immunol. Methods 196:17-32), peptides formulated as multivalent peptides; peptides for use in ballistic delivery systems, typically crystallized peptides, viral delivery vectors (Perkus, et al., p. 379 In: Kaufmann (ed. 1996) Concepts in vaccine development; Chakrabarti, et al. (1986) Nature 320:535; Hu, et al. (1986) Nature 320:537; Kieny, et al. (1986) AIDS Bio/Technology 4:790; Top, et al. (1971) J. Infect. Dis. 124:148; Chanda, et al. (1990) Virology 175:535), particles of viral or synthetic origin (see, e.g., Kofler, et al. (1996) J. Immunol. Methods 192:25; Eldridge, et al. (1993) Sem. Hematol. 30:16; Falo, et al. (1995) Nature Med. 7:649), adjuvants (Warren, et al. (1986) Annu. Rev. Immunol. 4:369; Gupta, et al. (1993) Vaccine 11:293), liposomes (Reddy, et al. (1992) J. Immunol. 148:1585; Rock (1996) Immunol. Today 17:131), or, naked or particle absorbed cDNA (Ulmer, et al. (1993) Science 259:1745; Robinson, et al. (1993) Vaccine 11:957; Shiver, et al., p. 423 In: Kaufmann (ed. 1996) Concepts in vaccine development; Cease and Berzofsky (1994) Annu. Rev. Immunol. 12:923 and Eldridge, et al. (1993) Sem. Hematol. 30:16). Toxin-targeted

delivery technologies, also known as receptor mediated targeting, such as those of Avant Immunotherapeutics, Inc. (Needham, Massachusetts) may also be used.

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Vaccine compositions often include adjuvants. Many adjuvants contain a substance designed to protect the antigen from rapid catabolism, such as aluminum hydroxide or mineral oil, and a stimulator of immune responses, such as lipid A, *Bortadella pertussis* or *Mycobacterium tuberculosis* derived proteins. Certain adjuvants are commercially available as, e.g., Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, MI); Merck Adjuvant 65 (Merck and Company, Inc., Rahway, NJ); AS-2 (SmithKline Beecham, Philadelphia, PA); aluminum salts such as aluminum hydroxide gel (alum) or aluminum phosphate; salts of calcium, iron or zinc; an insoluble suspension of acylated tyrosine; acylated sugars; cationically or anionically derivatized polysaccharides; polyphosphazenes; biodegradable microspheres; monophosphoryl lipid A and quil A. Cytokines, such as GM-CSF, interleukin-2, -7, -12, and other like growth factors, may also be used as adjuvants.

Vaccines can be administered as nucleic acid compositions wherein DNA or RNA encoding one or more of the polypeptides, or a fragment thereof, is administered to a patient. This approach is described, for instance, in Wolff, et. al. (1990) Science 247:1465 as well as U.S. Patent Nos. 5,580,859; 5,589,466; 5,804,566; 5,739,118; 5,736,524; 5,679,647; WO 98/04720; and in more detail below. Examples of DNA-based delivery technologies include "naked DNA", facilitated (bupivicaine, polymers, peptide-mediated) delivery, cationic lipid complexes, and particle-mediated ("gene gun") or pressure-mediated delivery (see, e.g., U.S. Patent No. 5,922,687).

For therapeutic or prophylactic immunization purposes, the peptides of the invention can be expressed by viral or bacterial vectors. Examples of expression vectors include attenuated viral hosts, such as vaccinia or fowlpox. This approach involves the use of vaccinia virus, e.g., as a vector to express nucleotide sequences that encode lung cancer polypeptides or polypeptide fragments. Upon introduction into a host, the recombinant vaccinia virus expresses the immunogenic peptide, and thereby elicits an immune response. Vaccinia vectors and methods useful in immunization protocols are described in, e.g., U.S. Patent No. 4,722,848. Another vector is BCG (Bacille Calmette Guerin). BCG vectors are described in Stover, et al. (1991) Nature 351:456-460. A wide variety of other vectors useful for therapeutic administration or immunization e.g., adeno and adeno-associated virus vectors, retroviral vectors, Salmonella typhi vectors, detoxified anthrax toxin vectors, and the

like, will be apparent to those skilled in the art from the description herein (see, e.g., Shata, et al. (2000) Mol Med Today 6:66-71; Shedlock, et al. (2000) J. Leukoc. Biol. 68:793-806; Hipp, et al. (2000) In Vivo 14:571-85).

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Methods for the use of genes as DNA vaccines are well known, and include placing a lung cancer gene or portion of a lung cancer gene under the control of a regulatable promoter or a tissue-specific promoter for expression in a lung cancer patient. The lung cancer gene used for DNA vaccines can encode full-length lung cancer proteins, but more preferably encodes portions of the lung cancer proteins including peptides derived from the lung cancer protein. In one embodiment, a patient is immunized with a DNA vaccine comprising a plurality of nucleotide sequences derived from a lung cancer gene. For example, lung cancer-associated genes or sequence encoding subfragments of a lung cancer protein are introduced into expression vectors and tested for their immunogenicity in the context of Class I MHC and an ability to generate cytotoxic T cell responses. This procedure provides for production of cytotoxic T cell responses against cells which present antigen, including intracellular epitopes.

In a preferred embodiment, DNA vaccines include a gene encoding an adjuvant molecule with the DNA vaccine. Such adjuvant molecules include cytokines that increase the immunogenic response to the lung cancer polypeptide encoded by the DNA vaccine. Additional or alternative adjuvants are available.

In another preferred embodiment lung cancer genes find use in generating animal models of lung cancer. When the lung cancer gene identified is repressed or diminished in metastatic tissue, gene therapy technology, e.g., wherein antisense or inhibitory RNA directed to the lung cancer gene will also diminish or repress expression of the gene. Animal models of lung cancer find use in screening for modulators of a lung cancer-associated sequence or modulators of lung cancer. Similarly, transgenic animal technology including gene knockout technology, e.g., as a result of homologous recombination with an appropriate gene targeting vector, will result in the absence or increased expression of the lung cancer protein. When desired, tissue-specific expression or knockout of the lung cancer protein may be necessary.

It is also possible that the lung cancer protein is overexpressed in lung cancer. As such, transgenic animals can be generated that overexpress the lung cancer protein.

Depending on the desired expression level, promoters of various strengths can be employed to express the transgene. Also, the number of copies of the integrated transgene can be determined and compared for a determination of the expression level of the transgene.

Animals generated by such methods will find use as animal models of lung cancer and are additionally useful in screening for modulators to treat lung cancer.

Kits for Use in Diagnostic and/or Prognostic Applications

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For use in diagnostic, research, and therapeutic applications suggested above, kits are also provided by the invention. In diagnostic and research applications such kits may include at least one of the following: assay reagents, buffers, lung cancer-specific nucleic acids or antibodies, hybridization probes and/or primers, antisense polynucleotides, ribozymes, RNAi, dominant negative lung cancer polypeptides or polynucleotides, small molecule inhibitors of lung cancer-associated sequences, etc. A therapeutic product may include sterile saline or another pharmaceutically acceptable emulsion and suspension base.

In addition, the kits may include instructional materials containing instructions (e.g., protocols) for the practice of the methods of this invention. While the instructional materials typically comprise written or printed materials they are not limited to such. A medium capable of storing such instructions and communicating them to an end user is contemplated by this invention. Such media include, but are not limited to electronic storage media (e.g., magnetic discs, tapes, cartridges, chips), optical media (e.g., CD ROM), and the like. Such media may include addresses to internet sites that provide such instructional materials.

The present invention also provides for kits for screening for modulators of lung cancer-associated sequences. Such kits can be prepared from readily available materials and reagents. For example, such kits can comprise one or more of the following materials: a lung cancer-associated polypeptide or polynucleotide, reaction tubes, and instructions for testing lung cancer-associated activity. Optionally, the kit contains biologically active lung cancer protein. A wide variety of kits and components can be prepared according to the present invention, depending upon the intended user of the kit and the particular needs of the user. Diagnosis would typically involve evaluation of a plurality of genes or products. The genes typically will be selected based on correlations with important parameters in disease which may be identified in historical or outcome data.

WO 02/086443 PCT/US02/12476 EXAMPLES

Example 1: Gene Chip Analysis

Molecular profiles of various normal and cancerous tissues were determined and analyzed using gene chips. RNA was isolated and gene chip analysis was performed as described (Glynne, et al. (2000) Nature 403:672-676; Zhao, et al. (2000) Genes Dev. 14:981-993).

Tables 1A and 1B were previously filed on April 18, 2001 in USSN 60/284,770 (18501-001500US) and on November 29, 2001 in USSN 60/334,370 (18501-001520US)

5	Table 1A					
9	Pkey	ExAccn	UnigenelD	Unigene Title	70% chron/90% NL	70% SQAD/90% NL
	100134	D13264	Hs.49	macrophage scavenger receptor 1	1.61	0.74
	100780 100971	HG3731-HT4001 J02874	Hs.83213	***Immunoglobulin Heavy Chain, Vdjrc Reg fatty acid binding protein 4; adipocyte	2.68 1.96	3.28 0.14
10		L05568	Hs.553	solute carrier family 6 (neurotransmitte	0.79	0.07
	101102	L07594	Hs.79059	transforming growth factor; beta recepto	2.55	1
		L15388	Hs.211569	G protein-coupled receptor kinase 5	0.88 0.89	0.27 0.26
	101277 101330	L38486 L43821	Hs.118223 Hs.80261	microfibrillar-associated protein 4 enhancer of filamentation 1 (cas-like do	0.59	0.29
15		L49169	Hs.75678	FBJ murine osteosarcoma viral oncogene h	1.15	0.41
	101345	L76380	Hs.152175	calcitonin receptor-like	0.81	0.31
	101678 101764	M62505 M80563	Hs.2161 Hs.81256	complement component 5 receptor 1 (C5a I S100 calcium-binding protein A4 (calcium	1.31 1,44	0.77 0.82
••	101771	M81750	Hs.153837	myeloid cell nuclear differentiation ant	0.96	0.45
20		M93221	Hs.75182	mannose receptor; C type 1	1.27	0.37
	102283 102363	U31384 U39447	Hs.83381 Hs.198241	guanine nucleotide blnding protein 11 amine oxldase; copper containing 3 (vasc	1.04 0.96	0.3 0.26
	102507	U52154	Hs.193044	potassium inwardly-rectifying channel; s	2.81	3.45
25	102698	U75272	Hs.1867	progastricsin (pepsinogen C)	0.95	0.23
25	103025	X54131	Hs.123641	protein tyrosine phosphatase; receptor t cadherin 5; VE-cadherin (vascular epithe	1.62 0.9	0.21 0.41
	103280 103496	X79981 Y09267	Hs.76206 Hs.132821	flavin containing monooxygenase 2	1,27	0.49
	103541	Z11697	Hs.79197	CD83 antigen (activated B lymphocytes; I	1.86	1
30			Hs.74034	caveolin 1; caveolae protein; 22kD	1.27	0.47
30	104212 104691	AB002298 AA011176	Hs.173035 Hs.37744	KIAA0300 protein ESTs	1.17 1.08	0.16 0.35
		AA035613	Hs.141883	ESTs	0.75	0.27
	104857	AA043219	Hs.19058	ESTs	2.6	3.3
35	104865 104989	AA045136 AA102098	Hs.22575 Hs.118615	ESTs ESTs	1.23 0.63	0.49 0.32
33	105729	AA292694	Hs.3807	ESTs; Weakly similar to PHOSPHOLEMMAN PR	0.86	0.34
	105847	AA398608	Hs.32241	ESTs	1.32	0.4
	105894 106490	AA400979 AA451861	Hs.25691 Hs.115537	calcitonin receptor-like receptor activi ESTs; Weakly similar to dipeptidase prec	0.78 1.2	0.28 0.47
40	106536	AA453997	Hs.23804	ESTs	0.82	0.15
• •	106605	AA457718	Hs.21103	Homo sapiens mRNA; cDNA DKFZp564B076 (fr	0.99	0.07
	106667	AA461086	Hs.16578	ESTS	1.17 1.46	0.4 0.43
	106773 106797	AA478109 AA478962	Hs.188833 Hs.169943	ESTs ESTs	1.18	0.43
45	106844	AA485055	Hs.158213	sperm associated antigen 6	0.98	0.51
	106870	AA487576	Hs.26530	serum deprivation response (phosphatidy)	1.05	0.14
	106954 107054	AA496980 AA600150	Hs.204038 Hs.14366	ESTs ESTs	1.25 1.11	0.33 0.4
	107292	T30407	Hs.4789	ESTs; Weakly similar to oxidative-stress	1.07	2.58
50	107994	AA036811	Hs.165030	ESTs	0.7	0.21
	107997 108041	AA037388 AA041552	Hs.82223 Hs.61957	Human DNA sequence from clone 141H5 on c ESTs	1.02 1.44	0.48 0.51
	108087	AA045709	Hs.40545	ESTs	1.98	1
55		AA074885	Hs.67726	macrophage receptor with collagenous str	1.52	0.72
22	108435 108480	AA078787 AA081093	Hs.194101 Hs.68055	ESTs .	2.53 1.56	1.53 0.48
	109252	AA194830	Hs.85944	ESTs	2.69	3.18
	109550	F01534	Hs.26981	ESTs	1.19	0.65
60	109613 109837	F03031	Hs.27519 Hs.29792	ESTs ESTs	1.01 0.81	0.29 0.15
00	109893	H04768	Hs.30484	ESTs	1.44	0.32
	109984	H09594	Hs.10299	ESTs	0.62	0.14
	110099 110837	H16568 N30796	Hs.23748 Hs.17424	ESTs ESTs; Weakly similar to semaphorin F [H.	1.01 1.1	0.28 0.22
65	111247	N69825	Hs.16762	Homo sapiens mRNA; cDNA DKFZp564B2062 (f	1.26	0.26
	111341	N80935	Hs.22483	ESTs	1.57	0.52
	111510	R07856 R25410	Hs.16355	ESTs ESTs	3.96 0.97	1 0.24
	111737 113195	T57112	Hs.9218	"'yc20g11.s1 Stratagene lung (#937210) .	1.22	0.35
70	113238	T62979	Hs.189813	ESTs	2.27	0.45
	113540	T90496	Hs.16757	ESTs	1.06 1.16	0.22 0.42
	113552 113606	T90889 T93093	Hs.16026 Hs.17125	ESTs ESTs	1.48	0.42
	113695	T96965	Hs.17948	ESTs	1.54	0.28
75	113945	W84753	Hs.37896	ESTs	1.79	0.72
	114251 114359	Z39898 Z41589	Hs.21948 Hs.153483	ESTs ESTs; Moderately similar to H1 chloride	1.95 1.42	0.25 0.13
	115230	AA278300	Hs.182980	ESTs	2.62	0.42
80	115279	AA279760	Hs.63671	ESTs	1.79	0.91
OU	115566 115965	AA398083 AA446661	Hs.43977 Hs.173233	ESTs ESTs	0.86 0.79	0.2 0.04
	116166	AA461556	Hs.202949	KIAA1102 protein	2.29	0.68
	116279	AA486073	Hs.57362	ESTs	2.27	0.78
	117023	H88157	Hs.41105	ESTs	1.36	0.16

	w	O 02/0864	43			
	117209	H99959	Hs.42768	ESTs	1.46	0.48
	118901	N90719	Hs.94445	ESTs	1.51	1
	118981	N93839	Hs.39288	ESTs	1.34	0.48
_	119073	R32894	Hs.45514	v-ets avian erythroblastosis virus E26 o	1.14	0.27
5	119221	R98105	11. 404	""yr30g11.s1 Soares fetal liver spleen	1.32 1	0.53 0.19
	119824 119861	W74536 W80715	Hs.184	advanced glycosylation end product-spect ESTs; Moderately similar to !!!! ALU SUB	1.83	0.15
	120041	W92775	Hs.59368	ESTs	1.23	0.55
	120132	Z38839	Hs.125019	ESTs; Highly similar to KIAA0886 protein	0.91	0.37
10	120467	AA251579	Hs.187628	ESTs	1.87	1.91
	121314	AA402799	Hs.182538	ESTs	1.3	0.31
	121643	AA417078	Hs.193767	ESTs	2.31	0.68
	121690	AA418074	Hs.110286	ESTS	1.47	0.51
15	122633	AA454080	Hs.34853	inhibitor of DNA binding 4; dominant neg	1.31	0.63
13	123978	C20653	Hs.170278	ESTs ESTs	1.52 0.93	0.32 0.35
	124214 124357	H58608 N22401	Hs.151323	""yw37g07.s1 Morton Fetal Cochlea Homo	1.29	1
	124438	N40188	Hs.102550	ESTs	1.36	0.7
	125167	W45560	Hs.102541	ESTs	1.46	0.69
20	125174	W51835	Hs.231082	EST	3.07	3.76
	125422	AA903229	Hs.153717	ESTs	1.34	0.3
	125561	A1417667	Hs.22978	ESTs	1.89	0.63
	125831	D60988		***HUM145B09B Clontech human fetal brain	0.94	0.36
25	127002	R35380	Hs.24979	ESTs	3.02	4.06 0.69
23	127307	AA369367	Hs.126712	ESTs; Weakly similar to pIL2 hypothetica	1.01 1.21	0.09
	127609 127959	AA622559 Al302471	Hs.150318 Hs.124292	ESTs ESTs	2.5	1
	128458	D52193	Hs.56340	ESTs	1.13	0.33
	128624	AA479209	Hs.102647	ESTs	1.45	. 0.58
30	128789	AA486567	Hs.105695	ESTs	1.1	0.34
	128798	AF014958	Hs.105938	chemokine (C-C motif) receptor-like 2	1.16	0.55
	128952	R51076	Hs.107361	ESTs; Highly similar to Rap2 interacting	2.04	2.4
	129057	X62466	Hs.214742	CDW52 antigen (CAMPATH-1 antigen)	1.77	0.73
25	129210	AA401654	Hs.202949	KIAA1102 protein	1,11	0.36
35	129240	W24360	Hs.237868	Interleukin 7 receptor	0.91	0.41
	129402 129565	T63781 X77777	Hs.198726	""yc21g01.s1 Stratagene lung (#937210) vasoactive intestinal peptide receptor 1	1.36 0.67	0.43 0.08
	129593	AA487015	Hs.98314	Homo sapiens mRNA; cDNA DKFZp586L0120 (f	1.3	0.42
	129626	AA447410	Hs.11712	ESTs; Weakly similar to IIII ALU SUBFAMI	1.28	0.46
40	129699	AA458578	Hs.12017	KIAA0439 protein; homolog of yeast ubiqu	1.58	1
	129898	N48595	Hs.13256	ESTs	1.13	0.53
	129958	L20591	Hs.1378	annexin A3	0.81	0.31
	130273	U59914	Hs.153863	MAD (mothers against decapentaplegic; Dr	0.59	0.22
15	130655	N92934	Hs.17409	cysteine-rich protein 1 (intestinal)	1.44	0.76
45	130657	T94452	Hs.201591	ESTS	0.96	0.42
	131061	N64328	Hs.22567	ESTs; Moderately similar to HYPOTHETICAL ESTs	1.51 0.97	0.45 0.37
	131066 131263	F09006 R38334	Hs.22588 Hs.24950	regulator of G-protein signalling 5	2.34	. 2.82
	131589	U52100	Hs.29191	epithelial membrane protein 2	1.2	0.62
50	131686	AA157428	Hs.30687	Grb2-associated binder 2	0.95	0.38
	131751	H18335	Hs.31562	ESTs	1.47	0.52
	132430	T23630	Hs.258675	EST	1.86	2.09
	132476	N67192	Hs.49476	Homo sapiens clone TUA8 Cri-du-chat regi	1.73	0.58
55	132836	F09557	Hs.57929	slit (Drosophila) homolog 3	0.91	0.29
23	133120	X64559 D45370	Hs.65424	tetranectin (plasminogen-binding protein	0.82 1.29	0.2 0.48
	133488 133565	H57056	Hs.74120 Hs.204831	adipose specific 2 ESTs	2.25	0.57
	133651	U97105	Hs.173381	dihydropyrimidinase-like 2	1.65	0.62
	133835	AA059489	Hs.76640	ESTs; Highly similar to RGC-32 [R.norveg	1.16	0.34
60	133978	W73859	Hs.78061	transcription factor 21	0.79	0.27
	133985	L34657	Hs.78146	platelet/endothelial cell adhesion molec	0.99	0.28
	134299	AA487558	Hs.8135	ESTs	1.02	0.46
	134300	U81984	Hs.166082	endothelial PAS domain protein 1	0.86	0.42
65	134323	AA028976	Hs.8175	Homo sapiens mRNA; cDNA DKFZp564M0763 (f	1.19	0.27
03	134343	D50683	Hs.82028 Hs.82921	transforming growth factor, beta recepto	1.21 1.28	0.67 1
	134417 134561	D87969 U76421	Hs.85302	solute carrier family 35 (CMP-sialic aci adenosine dearninase; RNA-specific; B1 (h	2.12	0.55
	134624	W67147	Hs.8700	deleted in liver cancer 1	2.35	2.74
	134696	H88354	Hs.8861	ESTs	1.35	0.33
70	134749	L10955	Hs.89485	carbonic anhydrase IV	0.89	0.2
	134786	L06139	Hs.89640	TEK tyrosine kinase; endothelial (venous	0.48	0.21
	134869	T3528B	Hs.90421	ESTs; Moderately similar to IIII ALU SUB	2.14	2.64
	135346	M21056	Hs.992	phospholipase A2; group IB (pancreas)	0.63	0.13
75	100113	D00591	Hs.84746	Chromosome condensation 1	1	2.15
13	100147	D13666	Hs.136348	Homo sapiens mRNA for osteoblast specifi	0.5 1.02	2 1.39
	100280 100335	D42085 D63391	Hs.155314 Hs.6793	KIAA0095 gene product platelet-activating factor acetythydrola	1.02	5.58
	100335	D78335	Hs.75939	Uridine monophosphate kinase	0.91	2.04
	100372	D79997	Hs.184339	KIAA0175 gene product	0.75	2.03
80	100486	HG1112-HT111		TIGR: ras-like protein TC4	1.09	1.93
	100559	HG2197-HT226		"collagen, type VII, aipha 1"	0.97	3.6
	100576	HG2290-HT238		"calcitonin/alpha-CGRP, alt. transcript	1	1
	100668	HG2981-HT393		*TIGR: CD44 (epican, alt. transcript 12	0.85	1.9
85	100906	HG4716-HT515		Guanosine 5'-Monophosphate Synthase	1.18	2.29 1.45
0,5	100930	HG721-HT4827		*TIGR: placental protein 14, endometrial	1	1.40

	W	U 02/080	443			
	100960	J00124	Hs.117729	keratin 14 (epidermolysis bullosa simple	0.84	26
	101031	J05070	Hs.151738	"Matrix metalloproteinase 9 (gelatinase	0.77	1.52
	101111	L08424	Hs.1619	Achaete-scute complex (Drosophila) homol	1	1
	101124	L10343	Hs.112341	"Protease inhibitor 3, skin-derived (SKA	0.62	2.67
5	101175	L18920	Hs.36980	"Melanoma antigen, family A, 2"	1	1
,				Ataxia-telangiectasia group D-associated	0.74	4.1
	101204	L24203	Hs.82237		0.85	2.51
	101431	M19888	Hs.1076	Small proline-rich protein 1B (comilin)		8.83
	101448	M21389	Hs.195850	keratin 5 (epidermolysis bullosa simplex	. 0.61	
4.0	101511	M27826	Hs.267319	Endogenous retroviral protease	1.03	1.13
10	101526	M29540	Hs.220529	Carcinoembryonic antigen-related cell ad	1.07	4.61
	101548	M31328	Hs.71642	*Guanine nucleotide binding protein (G p	0.97	1.13
	101625	M57293		"Human parathyroid hormone-related pepti	1	1
	101649	M60047	Hs.1690	Reparin-binding growth factor binding pr	1	2.7
	101724	M69225	Hs.620	bullous pemphigoid anligen 1 (230/240kD)	i	8.98
15					i	2.78
13	101748	M76482	Hs.1925	Desmoglein 3 (pemphigus vulgaris antigen		
	101759	M80244	Hs.184601	"Solute carrier family 7 (cationic amino	1.07	2.45
	101804	M86699	Hs.169840	TTK protein kinase	1	1
	101806	M86757	Hs.112408	S100 calcium-binding protein A7 (psorias	0.74	1.76
	101809	M86849		"Homo sapiens connexin 26 (GJB2) mRNA, c	1	7
20	101845	M93426	Hs.78867	*Protein tyrosine phosphatase, receptor-	1	1
	101851	M94250	Hs.82045	Midkine (neurite growth-promoting factor	1.13	2.6
	102083	U10323	Hs.75117	"Interleukin enhancer binding factor 2,	1.03	1.61
					0.94	3.62
	102154	U17760	Hs.75517	"Laminin, beta 3 (nicein (125kD), kalini		4.59
25	102193	U20758	Hs.313	secreted phosphoprotein 1 (osteopontin;	0.34	
25	102305	U33286	Hs.90073	chromosome segregation 1 (yeast homolog)	1.45	2.97
	102348	U37519	Hs.87539	Aldehyde dehydrogenase 8	0.52	2.25
	102581	U61145	Hs.77256	Enhancer of zeste (Drosophila) homolog 2	0.91	2.46
	102610	U65011	Hs.30743	Preferentially expressed antigen in meta	1	3.88
	102623	U66083	Hs.37110	"Melanoma antigen, family A, 9 (MAGE-9)"	1	1
30	102669	U71207	Hs.29279	Eyes absent (Drosophila) homolog 2	i	i
50		U74612		Forkhead box M1	1.06	2.77
	102696		Hs.239	Neurotensin	1.00	1
	102829	U91618	Hs.80962			
	102888	X04741	Hs.76118	Ubiquitin carboxyl-terminal esterase L1	1.13	2.59
	102913	X07696	Hs.80342	keratin 15	0.7	4.72
35	102915	X07820	Hs.2258	Matrix Metalloproteinase 10 (Stromolysin	1.15	3.35
	102963	X15943	Hs.37058	"Calcitonin/calcitonin-related polypepti	1	1
	103021	X53587	Hs.85266	"Integrin, beta 4"	1.38	2.34
	103036	X54925	Hs.83169	Matrix metalloprotease 1 (interstitial c	1	14.93
	103058	X57348	Hs.184510	Stratifin	1.25	4.17
40					1	1.72
1 0	103060	X57766	Hs.155324	matrix metalloproteinase 11 (stromelysin	1.16	7.38
	103119	X63629	Hs.2877	"Cadherin 3, P-cadherin (placental)"		
	103206	X72755	Hs.77367	monokine induced by gamma interferon	0.71	1.48
	103242	X76342	Hs.389	*Alcohol dehydrogenase 7 (class IV), mu	1	1
	103312	X82693	Hs.3185	*Lymphocyte antigen 6 complex, locus D;	0.92	1.28
45	103478	Y07755	Hs.38991	S100 calcium-binding protein A2	1.05	5.81
	103558	Z19574	Hs.2785	keratin 17	0.65	6.68
	103576	Z26317	Hs.2631	Desmoglein 2	0.79	1.73
			Hs.82128		1	3.93
	103587	Z29083 ,		5T4 Oncofetal antigen	0.71	7.23
50	103594	Z31560	Hs.816	"SRY (sex determining region Y)-box 2, p		
50	103768	AA089997		*ESTs, Highly similar to integral membra	0.99	1.8
	104158	AA454908	Hs.8127	KIAA0144 gene product	0.96	1.29
	104558	R56678	Hs.88959	Human DNA sequence from clone 967N21 on	1.23	7.23
	104689	AA010665		ESTs	0.96	2.11
	104733	AA019498	Hs.23071	ESTs	1.18	1.88
55	104906	AA055809	Hs.26802	Protein kinase domains containing protei	1.11	3,15
55			Hs.19322	ESTs; Weakly similar to !!!! ALU SUBFAMI	1.64	2.89
	104978 105012	AA088458 AA116036	Hs.9329	"Homo sapiens mRNA for fls353, complete	1.19	3.91
						4.63
	105175	AA186804	Hs.25740	ESTs; Wealdy similar to unknown [S.cerev	0.9	
C O	105263	AA227926	Hs.6682	ESTs	0.95	2.87
60 ·	105298	AA233459	Hs.26369	ESTs	1	1.13
	105312	AA233854	Hs.23348	S-phase kinase-associated protein 2 (p45	1.32	3.01
	105719	AA291644	Hs.36793	Hypothetical protein FLJ23188	1.28	2.31
	105743	AA293300	Hs.9598	ESTs	1	1
		AA411621	Hs.8895	ESTs; same as BFH6?	0.94	2.04
65	106231	AA429571	Hs.38002	KIAA1355 protein	1.04	1.5
05					1.26	2.26
	106540	AA454607	Hs.38114 Hs.105421	Hypothetical protein FLJ11100	1	2
	106575	AA456039		ESTs		
	106632	AA459897	Hs.11950	GPI-anchored metastasis-associated prote	0.87	1.32
7 0	106727	AA465342	Hs.34045	Hypothetical protein FLJ20764	0.87	1.59
70	106906	AA490237	Hs.222024	Transcription factor BMAL2 (cycle-like f	0.61	1.6
	107059	AA608545	Hs.23044	RAD51 (S. cerevislae) homolog (E coli Re	0.48	2.67
	107104	AA609786	Hs.15243	Nucleolar protein 1 (120kD)	1.01	1.44
	107151	AA621169	Hs.8687	ESTs; procollagen I-N proteinase	0.97	2.89
	107284	S74039	Hs.291904	Accessory proteins BAP31/BAP29	1.15	3.65
75		AA026418	Hs.91539	ESTs	0.72	3.44
15	107901				1	2.48
	107922	AA028028	Hs.61460	lg superfamily receptor LNIR precursor	-	
	107932	AA029317	Hs.18878	Hypothetical protein FLJ21620	1	1
	108695	AA121315	Hs.70823	KIAA1077 protein	0.91	3.53
00	108857	AA133250	Hs.62180	ESTs	1	1
80	108860	AA133334	Hs.129911	ESTs	0.73	7.3
-	108990	AA152296	Hs.72045	ESTs	1	1
	109166	AA179845	Hs.73625	*RAB6 interacting, kinesin-like (rabkine	i	4.55
	109424	AA227919	Hs.85962	Hyaluronan synthase 3	i	1.28
				Hypothetical protein DKFZp762H1311	1.42	2
85	109665	F05012	Hs.27027			2.16
ره	109970	H09281	Hs.13234	ESTs	1.13	۷. ان

	w	O 02/0864	43			
	110015	H10998	Hs.7164	A disintegrin and metalloproteinase doma	0.84	1.95
	110156	H18957	Hs.4213	ESTs	0.94	1.41
	110561	H59617	Hs.5199	HSPC150 protein similar to ubiquitin-con	0.91	3.18
_	111223	N68921	Hs.34806	ESTs; Weakly similar to neogenin (H.sapi	0.91	3.13 1.25
5	111345 111876	N89820 R38239	Hs.14559 Hs.293246	Hypothetical protein FLJ10540 *ESTs, Wealty similar to putative p150 [1 0.83	1.27
	111902	R39191	Hs.109445	KIAA1020 protein	0.91	0.91
	112244	R51309	Hs.70823	KIAA1077 protein	0.77	3.01
10	112973	T17271	11- 00004	"cDNA FLJ13308 fis, clone OVARC1001436,	1 0.55	1 1.03
10	112989 113047	T23482 T25867	Hs.89981 Hs.7549	*Diacylglycerol kinase, zeta (104kD)* ESTs	0.87	2
	113095	T40920	Hs.126733	ESTs	1	1
	113531	T90345	Hs.16740	Hypothetical protein FLJ11036	0.42	1.44
15	113970	W86748	Hs.8109	ESTs	1.17 0.86	1.73 0.82
15	114346 114407	Z41450 AA010188	Hs.130489 Hs.103305	*ATPase, aminophospholipid transporter-I ESTs	0.8	1.88
	114471	AA028074	Hs.104613	RP42 homolog	1.06	1.34
	114509	AA043551	Hs.101799	KIAA1350 protein	1.82	2.32
20	115060	AA253214	Hs.198249	"Gap junction protein, beta 5 (connexin	0.79 0.72	1.49 1.92
20	115091 115123	AA255900 AA256642	Hs.184523 Hs.236894	KIAA0965 protein "ESTs, High sim to LRP1_hu low density I	0.59	1.97
	115291	AA279943	Hs.122579	ESTs	1	1.25
	115506	AA292537	Hs.45207	Hypothetical protein KIAA1335	1.15	1.48
25	115522	AA331393	Hs.47378	ESTs	0.5 1	3.29 1
23	115536 115697	AA347193 AA411502	Hs.62180 Hs.63325	ESTs Homo sapiens type II membrane serine pro	i	6.53
	115909	AA436666	Hs.59761	ESTs	1	6.98
	115978	AA447522	Hs.69517	Differentially expressed in Fanconi anem	1	2.31
30	116028	AA452112	Hs.42644	thioredoxin-like	0.99 1.14	1.68 1.8
30	116107 116134	AA456968 AA460246	Hs.92030 Hs.50441	ESTs CGI-04 protein	1.11	1.86
	116157	AA461063	Hs.44298	Hypothetical protein	0.99	1.9
	116158	AA461187	Hs.61762	Hypoxia-inducible protein 2	0.44	0.86
35	116335	AA495830	Hs.87013	"Homo sapiens cDNA FLJ10238 fis, clone H	0.62 1.04	3.89 2.36
33	116483 117320	C14092 N23239	Hs.76118 Hs.211092	Ublquitin carboxyl-terminal esterase L1 LUNX protein; PLUNC(palate lung & nasal	0.51	0.64
	117557	N33920	Hs.44532	Diubiquitin	1.11	2.63
	117693	N40939	Hs.112110	PTD007 protein	0.98	1.79
40	117881	N50073	Hs.260622 Hs.48956	Butyrate-induced transcript 1 ESTs	1 0.67	1.43 2.86
40	118368 118566	N64339 N68558	Hs.42824	Hypothetical protein FLJ10718	1.21	0.83
	118695	N71781	Hs.50081	KIAA1199 see CVA7.doc	0.88	1.63
	119780	W72967	Hs.191381	ESTs: Weakly similar to hypothetical pro	1	1
45	119845	W79920	Hs.58561 Hs.132927	G protein-coupled receptor 87 "ESTs, Moderately similar to p53 regulat	1	1
73	120102 120104	W95428 W95477	Hs.180479	ESTs	Ò.69	3.07
	120486	AA253400	Hs.137569	Tumor protein 63 kDa with strong homolog	1.08	12.05
	120859	AA350158	Hs.1619	Achaete-scute complex (Drosophila) homol	1	1
50	120880 120948	- AA360240 AA397822	Hs.97019 Hs.104650	EST Hypothetical protein FLJ 10292	1.04	2.15
30	120983	AA398209	Hs.97587	EST	. 1	1
	121362	AA405500	Hs.97932	Chondromodulin I precursor	1	1.
	121369	AA405657	Hs.128791	CGI-09 protein "ESTs, Weakly similar to JM27 [H.sapiens	1	1.8 1
55	121791 123005	AA423978 AA479726	Hs.293317 Hs.105577	ESTs, veeled similar to JM27 (rt.sapiens	i	i
55	123044	AA481549	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro	0.95	1.88
	123160	AA488687	Hs.284235	ESTs	1.59	4.98
	123479	AA599469	Hs.135056	clone RP5-850E9 on chromosome 20 "ESTs, Weakly similar to PQ0109 Purkinje	1.19 1.03	1.64 1.14
60	123571 123829	AA608956 AA620697	Hs.112619 Hs.112208	XAGE-1 protein	1.39	2.2
•	124006	D60302	Hs.108977	ESTs	1	4.85
	124059	F13673	Hs.99769	ESTs	1.49 0.76	8.62 0.77
	124960 125218	T15386 W73561	Hs.194766 Hs.110024	Seizure related gene 6 (mouse)-like NADH:ubiquinone oxidoreductase MLRQ subu	1.33	1.77
65	125453	R06041	Hs.18048	"Melanoma antigen, family A, 10"	0.8	1.42
	125759	AA425587	Hs.82226	Glycoprotein (transmembrane) nmb	1.52	2.26
	125972	AA434562	Hs.35406	"ESTs, Highly similar to unnamed protein	1.05 1	2.48 1.95
	125994 126395	H55782 N70192	Hs.270799 Hs.278956	EST Hypothetical protein FLJ12929 .	i	1.35
70	126645	AJ167942	Hs.61635	STEAP1 (Homo sapiens BAC clone RG041D11	1	2.23
	127221	Al354332	Hs.72365	ESTs	0.73	3.27
	127479	AA513722	Hs.179729	collagen; type X; alpha 1 (Schmid metaph	0.51 1.8	1.94 3.16
	128192 128610	A1204246 L38608	Hs.10247	KIAA1085 protein activated leucocyte cell adhesion molecu	0.89	0.97
75	128777	U46006	Hs.10526	Cysteine and glycine-rich protein 2	1	1
	128924	AA234962	Hs.26557	Plakophilin 3	1.3	2.97
	129041	H58873	Hs.169902	"Solute carrier family 2 (facilitated gl	0.84 0.87	2.04 1.04
	129099 129404	H50398 AA172056	Hs.108660 Hs.111128	*ATP-binding cassette, sub-family C (CFT ESTs	1	1
80	129466	L42583		"Genbank Homo sapiens keratin 6 isoform	0.72	12.67
	129605	S72493	Hs.115947	Keratin 16 (focal non-epidermolytic palm	0.92	1.5 1.93
	129628 130023	U26727 X13461	Hs.1174 Hs.239600	"Cyclin-dependent kinase inhibitor 2A (m Calmodulin-like 3	0.85 0.84	1.22
	130023	X14850	Hs.147097	"H2A histone family, member X"	0.98	1.96
85	130385	AA126474	Hs.155223	stanniocalcin 2	1	1

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	130410	V01514	Hs.155421	Alpha-fetoprotein	0.63	0.63
	130441	U35835	Hs.301387	"Human DNA-PK mRNA, partial cds"	1.15	3.65
	130482	L32866	Hs.1578	Baculoviral IAP repeat-containing 5 (sur	1	1.88
-	130553	AA430032	Hs.252587	Pituitary tumor-transforming 1	0.92	1.98
5	130577	M35410	Hs.162	Insulin-like growth factor binding prote	1.17	4.7
	130627	L23808	Hs.1695	Matrix metalloproteinase 12 (macrophage	0.69	4.05
	130800	AA223386	Hs.19574	ESTs; Weakly similar to katanin p80 subu	1.13	241
	130939	AA598689	Hs.21400	ESTs	0.8	0.89
10	131046	X02530	Hs.2248	INTERFERON-GAMMA INDUCED PROTEIN PRECURS		1.15
10	131244	D38076	Hs.24763	RAN binding protein 1	1.13	1.85
	131877	J04088	Hs.156346	Topoisomerase (DNA) II alpha (170kD)	1	1
	131927	AA461549	Hs.34780	"Doublecortex; lissencephaly, X-linked (0.81	0.62
	131965	W90146	Hs.35962	ESTs	0.74	3.27
1.5	131978	D80008	Hs.36232	KIAA0186 gene product	1	1
15	132354	L05187	Hs.211913	Small proline-rich protein 1A	0.69	1.43
	132543	AA417152	Hs.5101	ESTs; Highly similar to protein regulati	0.79	4.27
	132632	N59764	Hs.5398	guanine-monophosphate synthetase	1	1.08
	132653	U31201	Hs.54451	"laminin gamma2 chain gene (LAMC2), exon	1	1
20	132659	Z75190	Hs.54481	*Low density lipoprotein receptor-relate	0.89	0.89
20	132710	W93726	Hs.55279	"Serine (or cysteine) proteinase inhibit	0.64	4.41
	132758	W52432	Hs.56105	*ESTs, Weakly similar to WDNM RAT WDNM1	1.55	2.08
	132767	L05188	Hs.231622	Small proline-rich protein 2B	0.83 0.55	1.66 0.55
	132816	M74542	Hs.575	Aldehyde dehydrogenase 3	1	3.53
25	132990	AA458761	Hs.18387	transcription factor AP-2 alpha (activat	1.16	2
23	133070	U69611	Hs.64311	"A disintegrin and metalloproteinase dom	1	2.7
	133282	U52960	Hs.286145	*SRB7 (suppressor of RNA polymerase B, y	0.95	1.42
	133317	AA215299	Hs.70830	U6 snRNA-associated Sm-like protein LSm7		2.55
	133370	AA156897	Hs.72157	Homo saplens mRNA; cDNA DKFZp564I1922	1.12	1.76
30	133391	X57579	Hs.727	H.sapiens activin beta-A subunit (exon 2	1.65 1.02	1.39
20	133832	H03387	Hs.241305	estrogen-responsive B box protein (EBBP)	1.02	1.55
	134032	Z81326	Hs.78589	"Serine (or cysteine) proteinase inhibit	0.95	1.53
	134168	AA398908	Hs.181634	The state of the s	1.36	2.48
	134218	AA227480	Hs.80205	Pim-2 oncogene ""collagen, type XI, alpha 1""	0.76	2.86
35	134405	R67275 X70683	Hs.82772 Hs.83484		1.89	3.78
55	134453 134470	X54942	Hs.83758	CDC28 protein kinase 2	1.82	4.11
	134645	U87459	Hs.167379		0.82	0.83
	134781	M17183	Hs.89626	Parathyroid hormone-like hormone	1	1
	135002	U19147	Hs.272484	Gantigen 6	i	i
40	100040	M97935	113.212707		0.92	1.25
10	101201	L22524	Hs.2256		2.92	8.5
	101664	M60752	Hs.121017		1	1
	102025	U03911	Hs.78934		0.8	1.61
	102023	U04898	Hs.2156		1	1
45	102221	U24576	110.2100		i	1
	102270	U30255	Hs.75888		1.08	1.43
	102339	U37022	Hs.95577		0.88	1.32
	102391	U41668	Hs.77494		1.07	1.58
	103000	X51956	Hs.146580		0.91	1.49
50	103395	X94754	Hs.119503		0.89	1.32
••	105638	AA281599	Hs.20418		0.91	1.25
	105726	AA292328	Hs.9754	activating transcription factor 5	0.94	1.48
	114841	AA234722	Hs.55408		0.78	1.56
	115206	AA262491	Hs.186572		1	1
55	115906	AA436616	Hs.82302		0.74	2.52
	119132	R49046	Hs.107911		1.1	1.51
	124163	H30539	Hs.189838	ESTs	1	1
	126487	AA482505	Hs.184601	solute carrier family 7 (cationic amino	1.01	1.46
	127141	AA307960	Hs.75478	KIAA0956 protein	0.85	1.4
60	128034	AA905754	Hs.75103	tyrosine 3-monooxygenase/tryptophan 5-mo	1	1.18
	128609	AA234365	Hs.102456	survival of motor neuron protein interac	1	1.5
	128895	R37753	Hs.106985	ESTs	1.7	2
	130199	Z48579	Hs.172028		1	1
	130524	U89995	Hs.159234		1	1
65	133000	U24152	Hs.62402	p21/Cdc42/Rac1-activated kinase 1 (yeast	1	1
	133658	M25756	Hs.75426	secretogranin II (chromogranin C)	1	1
	135047	AA460466	Hs.93597		1	1
	100053	M27830			0.88	1.53
~ ^	100114	D00596	Hs.82962		0.68	1.86
70	100128	D11094	Hs.61153		1.29	2.03
	100154	D14657	Hs.81892		0.71	4.26
	100161	D14694	Hs.77329		1.02	1.56
	100168	D14874	Hs.394 .		0.46	1.17
75	100187	D17793	Hs.78183		1	1
75	100188	D21063	Hs.57101		0.97	1.4
	100217	D26600	Hs.89545		1.13	1.9
•	100220	D28364			1.11	1.53
	100287	D43950	Hs.1600		1.13	2.09
00	100297	D49489	Hs.182429		0.92	1.78
80	100330	D55716	Hs.77152	minichromosome maintenance deficient (S.	1.07	1.61
	100355	D78129			0.96	1.87
	100364	D78586	Hs.154868	carbamoyl-phosphate synthetase 2; aspart	1.49	2.46
	100368	D79987	Hs.153479	extra spindle poles; S. cerevislae; homo	0.59	1.32
0.5	100398	D84557	Hs.155462	minichromosome maintenance deficient (m)	1.08	1.9
85	100438	D87448	Hs.91417	topoisomerase (DNA) II binding protein	1	2.15

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	100455	D87953	Hs.75789	N-myc downstream regulated	0.91	1.48
	100491	HG1153-HT115	3	Nucleoside Diphosphate Kinase Nm23-H2s	0.99	1.41
	100518	HG174-HT174		Desmoplakin I	1.28	3.17
_	100528	HG1828-HT185		""Nexin, Glia-Derived"	0.68	1.9
5	100661	HG2874-HT301		Ribosomal Protein L39 Homolog	1.1	5.44
	100667	HG2981-HT312		"Epican, Ait. Splice 11""	0.8	1.97
	100830	HG4074-HT434		Rad2	1.01	2.12 1.79
	101061	K03515	Hs.944	glucose phosphate isomerase splicing factor; arginine/serine-rich 3	0.91 1.23	1.75
10	101131 101162	L10838 L14595	Hs.167460 Hs.174203	solute cerrier family 1 (gluternate/neutr	1.35	2.73
10	101181	L19686	Hs.73798	macrophage migration inhibitory factor (1.03	1.78
	101183	L19779	Hs.795	H2A histone family; member O	0.57	1.3
	101216	L25876	Hs.84113	cyclin-dependent kinase inhibitor 3 (CDK	0.7	2.2
	101228	L27706	Hs.82916	chaperonin containing TCP1; subunit 6A (0.99	1.99
15	101233	L29008	Hs.878	sorbital dehydrogenase	0.82	2.11
	101247	L33801	Hs.78802	glycogen synthase kinase 3 beta	1.2	1.91
	101332	L47276	•	***Homo sapiens (cell line HL-6) alpha t	0.69	2.78
	101342	L76191	Hs.182018	interleukin-1 receptor-associated kinase	1.04	1.84
20	101396	M15796	Hs.78996	proliferating cell nuclear antigen	0.95	3.55
20	101423	M18391	Hs.89839	EphA1	1	1.5
	101445	M21259	Hs.1066	small nuclear ribonucleoprotein polypept	1.21	1.96
	101505	M27396	Hs.75692	asparagine synthetase	0.93	1.6
	101525	M29536	Hs.12163	eukaryotic translation initiation factor	1.19 0.96	1.93 1.42
25	101535 101607	M30448 M38690	Hs.251669 Hs.1244	casein kinase 2; beta polypeptide CD9 antigen (p24)	1.11	1.25
23	101624	M55998	115,1244	""*Human alpha-1 collagen type I gene, 3	1.17	1.98
	101758	M77836	Hs.79217	pyrroline-5-carboxylate reductase 1	1.77	3.45
	101839	M93036	Hs.692	membrane component; chromosomal 4; surfa	0.71	1.45
	101853	M94362	Hs.76084	lamin B2	0.84	1.19
30	101977	S83364		""putative Rab5-interacting protein (cl	0.89	1.9
	101992	U01038	Hs.77597	polo (Drosophia)-like kinase	0.66	1.46
	102009	U02680	Hs.82643	protein tyrosine kinase 9	1.23	3.35
	102012	U03057	Hs.118400	singed (Drosophila)-like (sea urchin fas	0.85	1.88
25	102039	U05861	Hs.201967	aldo-keto reductase family 1; member C1	0.93	2.32
35	102123	U14518	Hs.1594	centromere protein A (17kD)	1	4.28
	102130	U15009	Hs.1575	small nuclear ribonucleoprotein D3 polyp	0.89	1.42 2.95
	102148	U16954	Hs.75823	ALL1-fused gene from chromosome 1q	0.8 1.01	2.55 1.34
	102210 102220	U23028 U24389	Hs.2437 Hs.65436	eukaryotic translation initiation factor lysyl oxidase-like 1	1.15	2.34
40	102220	U28386	Hs.159557	karyopherin alpha 2 (RAG cohort 1; Impor	1.14	2.69
40	102330	U35451	Hs.77254	chromobox homolog 1 (Drosophila HP1 beta	1.05	1.7
	102423	U44754	Hs.179312	small nuclear RNA activating complex; po	1,14	2.99
	102455	U48705	Hs.75562	discoidin domain receptor family; member	1.05	2.01
	102499	U51478	Hs.76941	ATPase; Na+/K+ transporting; beta 3 poly	1.27	1.92
45	102522	U53347	Hs.183556	solute carrier family 1 (neutral amino a	0.84	1.31
	102590	U62136		""Homo sapiens enterocyte differentiati	1.11	1.6
	102676	U72514	Hs.12045	putative protein	1.04	2.17
	102687	U73379	Hs.93002	ubiquitin carrier protein E2-C	0.86	2.28
50	102704	U76638	Hs.54089	BRCA1 associated RING domain 1	1.12	1.63 1.39
50	102781	U83843	Un 61706	****Human HIV-1 Nef interacting protein (0.9 0.98	2.16
	102784 102827	U85658 U91327	Hs.61796 Hs.6456	transcription factor AP-2 gamma (activat chaperonin containing TCP1; subunit 2 (b	0.96	1.62
	102935	X13482	Hs.80506	small nuclear ribonucleoprotein polypept	1.21	4.2
	102972	X16662	Hs.87268	annexin A8	1.25	2.32
55	102983	X17620	Hs.118638	non-metastatic cells 1; protein (NM23A)	1.03	1.83
	103023	X53793	Hs.117950	multifunctional polypeptide similar to S	1.58	5.44
	103038	X54941	Hs.77550	CDC28 protein kinase 1	1.32	3.79
	103075	X59543	Hs.2934	ribonucleotide reductase M1 polypeptide	1,11	2.58
C O	103168	X68314	Hs.2704	glutathione peroxidase 2 (gastrointestin	0.75	3.05
60	103185	X69910	Hs.74368	transmembrane protein (63kD); endoplasmi	1.01	1.97
	103212	X73874	Hs.2393	phosphorylase kinase; alpha 1 (muscle)	0.95 0.97	1.72 1.77
	103223	X74801	Hs.1708 Hs.3155	chaperonin containing TCP1; subunit 3 (g	0.97 1	1.77
	103260 103262	X78416 X78565	Hs.204133	casein; alpha hexabrachion (tenascin C; cytotactin)	1.23	3.09
65	103330	X85373	Hs.77496	small nuclear ribonucleoprotein polypept	1.12	2.25
00	103364	X90872	Hs.75854	SULT1C sulfotransferase	2.85	4.62
	103375	X91868	Hs.54416	sine oculis homeobox (Drosophila) homolo	1	2.48
	103391	X94453	Hs.114366	pyrroline-5-carboxylate synthetase (glut	1	1.53
	103404	X95586	Hs.78596	proteasome (prosome; macropain) subunit;	0.92	1.53
70	103437	X98260	Hs.82254	M-phase phosphoprotein 11	0.92	1.54
	103448	X99133	Hs.204238	lipocalin 2 (oncogene 24p3)	0.55	0.96
	103605	Z35402	Hs.194657	cadherin 1; E-cadherin (epithelial)	1.32	2.51
	103646	Z68228	Hs.2340	junction plakoglobin	0.88	1.28 2.98
75	103658	Z74615	Hs.172928	collagen; type I; alpha 1 ESTs; Weakly similar to R07G3.8 [C.elega	1.06 1.88	2.98 4.66
, 5	103774 104261	AA092898 AF008442	Hs.9291B Hs.5409	RNA polymerase I subunit	0.87	2.17
	104261	C02193	Hs.85222	ESTs; Weakly similar to R27090_2 [H.sapi	1.4	2.49
	104270	C16281	Hs.75478	KIAA0956 protein	1.15	1.68
	104434	L02870	Hs.1640	collagen; type VII; alpha 1 (epidermolys	1.04	1.49
80	104453	M19169	Hs.123114	cystalin SN	0.38	0.76
	104611	R98280	Hs.125845	ributose-5-phosphate-3-epimerase	1.08	2.25
	104758	AA024661	Hs.7010	ESTs; Weakly similar to ACYL-COA DEHYDRO	1.14	1.65
	105114	AA156532	Hs.11801	adenosine A2b receptor pseudogene	0.91	1.38
85	105132	AA159501	Hs.247280	HBV associated factor	1.08	1.7
O)	105174	AA186613	Hs.34744	ESTs	0.95	2.05

	W	O 02/086	5443			
	105280	AA232215	Hs.14600	ESTs	1	1.4
	105344	AA235303	Hs.8645	ESTs	0.72	2.02
	105516	AA257971	Hs.21214	EST8	1.35	3.56
_	105621	AA280865	Hs.6375	Homo sapiens mRNA; cDNA DKFZp564K0222 (f	1.23	1.82
5	105698	AA287393	Hs.15202	ESTs; Weakly similar to oligodendrocyte-	0.98	1.28
	105705	AA290767	Hs.101282	Homo sepiens mRNA; cDNA DKFZp434B102 (fr	0.92	1.32
	105724	AA292098	Hs.22934	ESTs; Weakly similar to ZINC FINGER PROT	0.99	1.41
	105782	AA350215	Hs.21580	ESTs	1	1
10	105799	AA372018	Hs.24743	ESTs	1.08	1.78
10	105807	AA393803	Hs.16869	ESTs; Moderately similar to COLLAGEN ALP	0.95	1.34
	105891	AA400768	Hs.26662	ESTs; Weakly similar to tumor necrosis f	0.87	2.25
	105936	AA404338		ESTs	1.14	1.46
	106069	AA417741	Hs.29899	ESTs; Weakly similar to ZINC FINGER PROT	1	1 1.44
16	106103	AA421104	Hs.12094	ESTs	1.04	2.11
15	106140	AA424524	Hs.14912	KIAA0286 protein	1.23	
	106149	AA424881	Hs.256301	ESTs	0.83 0.77	1.48 2.05
	106154	AA425304	Hs.6994	ESTs	0.74	2.23
	106182	AA426609	Hs.10862	ESTs Madaustah alamba ta matamidis s	0.74	1.99
20	106220	AA428582	Hs.32196	ESTs; Moderately similar to metargidin p	0.99	1.54
20	106228	AA429290	Hs.17719	ESTS	0.95	2.09
	106318	AA436570	Hs.9605	pre-mRNA cleavage factor Im (25kD)	0.98	2.66
	106341	AA441798	Hs.5243	ESTs; Moderately similar to pIL2 hypothe	0.95	1.93
	106432	AA448850	Hs.17138	ESTS Home capiens mBNA: aDNA DKE7a564C053 (fr	1	1
25	106474	AA450212	Hs.42484	Homo sapiens mRNA; cDNA DKFZp564C053 (fr IGF-II mRNA-binding protein 2	1.4	2.29
23	106483	AA451676	Hs.30299	ESTs; Moderately similar to non-function	i	1.82
	106599	AA457235	Hs.12842		1.49	2.78
	106611	AA458904	Hs.26267 Hs.3784	ESTs; Weakly similar to torsinA [H.sapie ESTs; Highly similar to phosphoserine am	1	1.4
	106654	AA460449	Hs.21143	ESTs; Weakly similar to prospression and	1.11	1.49
30	107076	AA609145	Hs.27693	ESTs; Highly similar to CGI-124 protein	1	1.03
50	107115 107129	AA610108 AA620553	Hs.4756	flap structure-specific endonuclease 1	1,13	3.63
	107159	AA621340	Hs.10600	ESTs; Weakly similar to ORF YKR081c [S.c	1.05	2.09
	107444	W28391	Hs.5181	proliferation-associated 2G4; 38kD	1.18	1.9
	107481	W58247	Hs.27437	Homo sapiens kinesin superfamily motor K	0.99	2.74
35	107516	X56597	Hs.99853	fibrillarin	0.94	1.77
55	107529	Y12065	Hs.5092	nucleolar protein (KKE/D repeat)	1.05	2.29
	107531	Y13936	Hs.17883	protein phosphatase 1G (formerly 2C); ma	1.06	1.62
	107801	AA019433	Hs.173100	ESTs	1.03	1.4
	107957	AA031948	Hs.57548	ESTs	0.95	1.46
40	108565	AA085342	Hs.1526	ATPase; Ca++ transporting; cardiac muscl	0.59	1.35
	108780	AA128561	Hs.117938	collagen; type XVII; alpha 1	1	7.63
	108828	AA131584	Hs.71435	DKFZP564O0463 protein	1.33	2.56
	109060	AA160879	Hs.241551	chloride channel; calcium activated; fam	0.67	1.42
	109112	AA169379	Hs.72865	ESTs	1.03	2.31
45	109344	AA213696	Hs.86559	poly(A)-binding protein-like 1	0.97	1.55
	109412	AA227145	Hs.209473	ESTs; Weakly similar to REGULATOR OF MIT	0.76	1.87
	110780	N23174	Hs.22891	solute carrier family 7 (cationic amino	0.9	0.95
	110958	N50550	Hs.24587	signal transduction protein (SH3 contain	1.17	2.26
	111018	N54067	Hs.3628	mitogen-activated protein kinase kinase	1.21	1.85
50	111337	N79612	Hs.16607	ESTs; Highly similar to Myosin heavy cha	1	1.45
	112305	R54822	Hs.26244	ESTs	1	1
	112401	R61279	Hs.237536	ESTs; Weakly similar to F25B5.3 [C.elega	1.24	1.64
	112853	T02843	Hs.4351	EST	1.56	1.96
	112869	T03313	Hs.4747	dyskeratosis congenita 1; dyskertn	1.03	1.57
55	112992	T23513	Hs.7147	ESTs .	1	1
	113048	T25895	Hs.184008	ESTs; Weakly similar to RNA-binding prot	1.37	2.26
	113063	T32438	Hs.5027	ESTs	1	· 1_
	113179	T55182	Hs.152571	ESTs; Highly similar to IGF-II mRNA-bind	1.33	2.7
C 0	113573	T91166	Hs.15990	ESTs	0.76	1.47
60	113811	W44928	Hs.4878	ESTs	0.79	1.51
	114086	Z38266	Hs.12770	Homo sapiens PAC clone DJ0777O23 from 7p	0.9	1.34
	114587	AA070827	Hs.180320	ESTs; Weakly similar to GOLGI 4-TRANSMEM	1.02	1.76
	114846	AA234929	Hs.44343	ESTs	1.32	2.36
65	114964	AA243873	Hs.82184	ring finger protein 3	1.1	1.84
65	115047	AA252627	Hs.22554	homeo box B5	1.01	2.36
	115166	AA258409	Hs.198907	myelin protein zero-like 1	1.05	2.31
	115167	AA258421	Hs.43728	hypothetical protein	1.52	2.52
	115239	AA278650	Hs.73291	ESTs; Weakly similar to similar to the b	0.7	2.57 2.12
70	115278	AA279757	Hs.67466	ESTs; Weakly similar to BACN32G11.d [D.m	1.14 0.82	4.67
70	115652	AA405098	Hs.38178	ESTS	1.2	1.98
	115875	AA433943	Hs.43946	ESTs; Weakly similar to Weak similarity	0.96	1.31
	116004	AA449122	Hs.76086	ESTs; Highly similar to small zinc finge	0.97	1.55
	116121	AA459254	Hs.48855	ESTs	1.08	2.73
75	116129	AA459956	Hs.49163	ESTs; Highly similar to putative ribonuc	0.8	1.57
, 5	116190	AA464963	Hs.67776	EST6	1.37	2.65
	116312	AA490494 E13770	Hs.65403	ESTs ESTs	0.92	1.8
	116732	F13779 N35020	Hs.165909 He 44685	ESTs; Weakly similar to GOLIATH PROTEIN	1.15	1.84
	117602 117950	N51394	Hs.44685 Hs.75478	KIAA0956 protein	1.04	2.36
80	117992	N52000	Hs.172089	Homo saplens mRNA; cDNA DKFZp586B0222 (f	0.62	1.29
-	118785	N75386	Hs.111867	GLI-Kruppel family member GLI2	1	1
	119717	W69134	Hs.57987	ESTs	i	1.4
	119814	W74069	Hs.58350	ESTs	0.78	1.77
	120128	Z38499	Hs.91448	MKP-1 like protein tyrosine phosphatase	0.86	1.46
85	120242	Z98443	Hs.86366	ESTs	0.83	2.01
				· -		•

	w	O 02/086	5443			
	120483	AA252994	Hs.1578	apoptosis inhibitor 4 (survivin)	0.74	1.64
	121054	AA398604	Hs.97387	ESTs	1.05	1.93
	121326	AA404246	Hs.97031	ESTs; Weakly similar to Similar to phyto	0.98	1.3
5	121376	AA405699	Hs.166232	ESTs; Moderately similar to SODIUM- AND	0.91	1.83 1.59
)	121457 121780	AA411448 AA422086	Hs.208985 Hs.124660	ESTs ESTs	0.91 0.46	0.55
	121781	AA422150	Hs.98370	cytochrome P540 family member predicted	1.07	1.54
	121844	AA425732	Hs.98485	gap junction protein; beta 2; 26kD (conn	0.94	1.4
10	122059	AA431737	Hs.98749	EST	1.93	2.33
10	122338	AA443311	Hs.98998	ESTs	1	1
	122354	AA443772	Hs.186692	ESTs . Wealth similar to MP (L) contact	0.88 2.28	1.39 2.93
	122591 122790	AA453265 AA460156	Hs.99311 Hs.99556	ESTs; Weakly similar to MRJ [H.saplens] ESTs	0.88	1.3
	123398	AA521265	Hs.105514	ESTs ·	1	1.93
15	123518	AA608531	Hs.170313	ESTs	1	1
	123673	AA609471	Hs.112712	ESTs	1	1.15
	124000	D57317	Hs.74861	activated RNA polymerase II transcriptio	0.74 0.67	1.12 1.1
	124367 124447	N24006 N48000	Hs.99348 Hs.140945	distal-less homeo box 5 Homo sapiens mRNA; cDNA DKFZp586L141 (fr	1.19	1.7
20	125756	W25498	Hs.81634	ATP synthase; H+ transporting; mitochond	0.93	1.59
	125769	Al382972	Hs.82128	5T4 oncofetal trophoblast glycoprotein	1.65	6.76
	125852	H09290	Hs.76550	Homo sapiens mRNA; cDNA DKFZp564B1264 (f	0.72	2.26
	125924	AA526849	Hs.82109	syndecan 1	1.22 1.36	2.25 1.63
25	126037 126214	M85772 N29455	Hs.6066 Hs.74316	KIAA1112 protein desmoplakin (DPI; DPII)	1.93	3.55
23	126414	N78770	Hs.223439	ESTs	1.21	1.66
	126737	AA488132	Hs.62741	ESTs .	1	1
	126743	AA179253	Hs.172182	poly(A)-binding protein; cytoplasmic 1	1.3	2.16
30	126926	AA179546	Hs.832 Hs.170311	ESTs; Highly similar to INTEGRIN BETA-8 heterogeneous nuclear ribonucleoprotein	2,53 1.57	2.8 2.12
30 .	127432 128218	AA501734 H02682	Hs.99189	ESTs; Moderately similar to recombinatio	1.24	2.09
	128527	M31523	Hs.101047	transcription factor 3 (E2A immunoglobul	1.08	1.78
	128568	X60673	Hs.247568	adenylate kinase 3	1.23	3.48
25	128584	M11433	Hs.101850	retinol-binding protein 1; cellular	0.87	2.42
35	128628 128691	C14037 W27939	Hs.251978 Hs.103834	EST ESTs	1.22 1.1	1.9 1.73
	128714	V00599	Hs.179661	Homo sapiens clone 24703 beta-tubulin mR	0.92	1.17
	128733	AA328993	Hs.104558	EST8	1.34	1.94
40	128781	X85372	Hs.105465	small nuclear ribonucleoprotein polypept	0.9	1.34
40	129052 129095	AA496297 L12350	Hs.182740 Hs.108623	ribosomal protein S11 thrombospondin 2	2.59 1.04	3.19 3.2
	129093	AA435665	Hs.109706	ESTs; Moderalely similar to HN1 [M.muscu	0.95	1.61
	129665	M88458	Hs.118778	KDEL (Lys-Asp-Glu-Leu) endoplasmic retic	1.28	2.63
	129703	AA401348	Hs.179999	ESTs	0.97	1.63
45	129720	AA476582 .	Hs.12152	ESTs; Moderately similar to SIGNAL RECOG GDP dissociation inhibitor 2	1.09 0.74	1.79 1.68
	129850 129896	N20593 AA043021	Hs.56845 Hs.13225	UDP-Gal:belaGlcNAc beta 1;4- galactosylt	1.43	4.19
	130069	AA055896	Hs.146428	collagen; type V; alpha 1	1.17	1.98
~ 0	130405	H88359	Hs.155396	nuclear factor (erythroid-derived 2)-lik	1.26	1.79
50	130541	X05608	Hs.211584	neurofilament; light polypeptide (68kD)	1 1.07	1 1.66
	130599 130867	M91670 J04093	Hs.174070 Hs.2056	ubiquitin carrier protein UDP glycosyltransferase 1	1.07	4.8
	131009	AA063596	Hs.22142	ESTs; Weakly similar to NADH-CYTOCHROME	0.93	1.05
	131028	U20240	Hs.2227	CCAAT/enhancer binding protein (C/EBP);	1	1.23
55	131083	U66661	Hs.22785	gamma-aminobutyric acid (GABA) A recepto	1.1	1.8
	131091	T35341	Hs.22880	ESTs; Highly similar to dipeptidyl pepti ESTs; Highly similar to HSPC038 protein	1.28 1.43	1.98 2.06
	131144 131148	C14412 C00038	Hs.23528 Hs.23579	ESTs Figury Surinal to Horocoo protein	0.88	3.38
	131164	Y00503	Hs.182265	keratin 19	1.19	2.77
60	131185	M25753	Hs.23960	cyclin B1	0.86	3.84
*	131219	C00476	Hs.24395	small inducible cytokine subfamily B (Cy	0.66	2.96
	131454	AA455896 L11066	Hs.2699 Hs.3069	glypican 1 heat shock 70kD protein 9B (mortalin-2)	0.99 1	1.54 1.18
	131687 131689	AA599653	Hs.30696	transcription factor-like 5 (basic helix	i	1.95
65	131692	D50914	Hs.30736	KIAA0124 protein	1.55	2.39
	131786	AA135554	Hs.32125	ESTs	1	1.33
1	131843	AA195893	Hs.184062	ESTs; Moderately similar to putative Rab	0.83 1.08	1.63 2.2
	131860 131884	U02082 H90124	Hs.334 Hs.3463	Oncogene TIM ribosomal protein S23	1.23	1.24
70	131903	AA481723	Hs.3436	deleted in oral cancer (mouse; homolog)	0.91	1.18
	131945	M87339	Hs.35120	replication factor C (activator 1) 4 (37	1	2.8
	131958	AA093998	Hs.3566	ESTs; Highly similar to phosphorylation	0.87	1.36
	131964	W42508	Hs.3593	ESTs v-Ha-ras Harvey rat sarcoma viral oncoge	1 1.12	1.25 1.43
75	132001 132040	J00277 AA146843	Hs.37003 Hs.172894	BH3 Interacting domain death agonist	1	1.55
	132065	D82226	Hs.211594	proteasome (prosome; macropain) 26S subu	0.89	1.27
	132109	AA599801	Hs.40098	ESTs	1	1.05
	132112	AA150661	Hs.40154	jumonji (mouse) homolog	0.99	1.44
80	132123	AA447123	Hs.250705	ESTs ESTs	1.06 1.08	2.46 2.46
00	132162 132180	H89551 AA405569	Hs.41241 Hs.418	fibroblast activation protein; alpha; se	1.02	4.56
	132309	AA460917	Hs.2780	jun D proto-oncogene	1.16	1.8
	132371	AA235448	Hs.46677	ESTs	0.8	1.26
85	132618	AA253330	Hs.5344	adaptor-related protein complex 1; gamma	0.5 1,21	1.49 1.81
0.5	132736	U68019	Hs.211578	MAD (mothers against decapentaplegic; Dr	1,6.1	1.01

	w	O 02/0864	143				PCT/US02/12476
	132771	AA488432	Hs.56407	phosphoserine phosphatase	1	1.3	
	132833	U78525	Hs,57783	eukaryotic translation initiation factor	0.91	1.43	
	132922	T23641	Hs.6066	KIAA1112 protein	1.16	1.53	
	132959	AA028103	Hs.61472	ESTs; Wealdy similar to unknown [S.cerev	1.02	1.88	
5	132994	AA505133	Hs.7594	solute carrier family 2 (facilitated glu	0.72	2.97	
					0.88	1.34	
	133005	C21400	Hs.103329	KIAA0970 protein	0.93	1.23	
	133065	X62535	Hs.172690	diacytglycerol kinase; alpha (80kD)	0.53 1.14	1.76	
	133083	N70633	Hs.6456	chaperonin containing TCP1; subunit 2 (b			
10	133086	L17131	Hs.139800	high-mobility group (nonhistone chromoso	0.97	1.43	
10	133134	T89703	Hs.65648	RNA binding motif protein 8	1.1	1.8	
	133195	AA350744	Hs.181409	KIAA1007 protein	2.29	2.69	
	133313	AA249427	Hs.70704	ESTs	1.07	1.68	
	133331	T62039	Hs.158675	ribosomal protein L14	0.85	1.18	•
	133438	D13370	Hs.73722	APEX nuclease (multifunctional DNA repai	0.91	1.45	
15	133445	T99303	Hs.73797	guanine nucleotide binding protein (G pr	0.94	1.68	
	133483	X52426	Hs.74070	keratin 13	0.85	1.14	
	133492	L40397	Hs.74137	transmembrane trafficking protein	1.1	1.69	
	133504	W95070	Hs.74316	desmoplakin (DPI; DPII)	0.7	6.21	
	133517	X52947	Hs.74471	gap junction protein; alpha 1; 43kD (con	0.95	1.3	
20	133540	D78151	Hs.74619	proteasome (prosome; macropain) 26S subu	0.91	1.25	
20	133594	L07758	Hs.172589	nuclear phosphoprotein similar to S. cer	0.84	1.29	
	133627	U09587	Hs.75280	glycyl-tRNA synthetase	1.09	1.99	
	133671	T25747	Hs.75471	zinc finger protein 146	1.02	1.5	
	133859	U86782	Hs.178761	26S proteasome-associated pad1 homolog	1.11	3.33	
25		F09315	Hs.170290	discs; large (Drosophila) homolog 5	1.84	6.7	
23	133865				1.15	1.86	
	133913	W84712	Hs.7753	calumenin	1.3	1.91	
	133963	L34587	Hs.184693	transcription elongation factor B (SIII)	1.3 1.3	1.99	
	133982	U47621	Hs.207251	nucleolar autoantigen (55kD) similar to	0.72	1.65	
20	134100	L07540	Hs.171075	replication factor C (activator 1) 5 (36		1.62	
30	134110	U41060	Hs.79136	LIV-1 protein; estrogen regulated	1.04		
	134158	U15174	Hs.79428	BCL2/adenovirus E1B 19kD-interacting pro	1	1.55	
	134161	U97188	Hs.79440	IGF-II mRNA-binding protein 3	0.82	1.95	
	134193	F09570	Hs.7980	ESTs	0.98	1.48	
~ -	134367	X54199	Hs.82285	phosphoribosylglycinamide formyltransfer	1	2.8	
35	134402	U25165	Hs.82712	fragile X mental retardation; autosomal	1.26	2	
	134457	D86963	Hs.174044	dishevelled 3 (homologous to Drosophila	1	1.47	
	134469	X17567	Hs.83753	small nuclear ribonucleoprotein polypept	0.94	1.57	•
	134498	M63180	Hs.84131	threonyl-tRNA synthetase	1.2	2.64	
	134501	W84870	Hs.211568	eukaryotic translation initiation factor	0.84	1.36	
40	134507	M63488	Hs.84318	replication protein A1 (70kD)	1.7	2.93	
• •	134548	U41515	Hs.85215	Deleted in split-hand/split-foot 1 regio	1.46	2.73	
,	134599	X99226	Hs.86297	Fanconi anemia; complementation group A	1.36	2.22	
	134692	R73567	Hs.8850	a disintegrin and metalloproteinase doma	0.77	1.64	
	134693	N70361	Hs.8854	ESTs	1.09	1.82	
45	134806	Z49099	Hs.89718	spermine synthase	0.98	1.35	
10	134821	Z34974	Hs.198382	plakophilin 1 (ectodermal dysplasia/skin	0.99	1.4	
	134864	Y08999	Hs.90370	actin related protein 2/3 complex; subun	0.95	1.42	
	134914	U29615	Hs.91093	chitinase 1 (chitotriosidase)	1.16	1.29	
	134953	L10678	Hs.91747	profilin 2	0.95	1.76	
50	134993	AA282343	Hs.9242	purine-rich element binding protein B	0.9B	1.73	
20		C15324	Hs.93668	ESTs	1.35	2.11	
	135051		115.53000	Human desmocollin-2 mRNA; 3' UTR	0.86	1.16	
	135158	U51711		Human upanocomiez mana, o OTA	0.00	1.10	

Table 1B shows the accession numbers for those pkeys in Table 1A lacking unigenelD's. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the Accession column.

60	CAT number:	Unique Eos probeset identifier number Gene cluster number
	Accession:	Genbank accession numbers

	Pkey	CAT	Accessions
65	100661 100667	23182_1 26401_3	BE623001 L05096 AA383604 AW966416 N53295 AA460213 AW571519 AA603655 L05424 X56794 S66400 X55150 W60071 AW351820 X55938 M83326 BE005289 BE070059 M83324 BE005248 BE069717 BE181648 BE069700 AW606203 BE069721 AW382138 AW803776 BE463954 BE005334 BE005274 T27386 AA932714 AA972695 AW377728 AI632506 T250044
70			A1783934 AW377727 BE163715 AL047291 AA279047 AA523003 BE008048 BE440141 W23614 BE090519 BE092193 N29181 N20368 N44153 BE546944 T692211 AW377441 AA907406 H50799 AW051416 A1420712 BE620922 A1279161 AA992549 W47198 BE005241 A1342696 H50700 A1969974 A1863855 AA374490 AW130675* A1950633 AA146667 H99482 X55150 BE005414 BE005339 N28294 A1673068 A1887890 AW804171 A1675961 AW804172 AA778841 AL048050 A127757 A1095568 AW204955 AW68978 W31898 A1052595 A1278771 BE464018 A1081503 A1824196 AA513211 AA411062 AW084376 N48752 AA703209 N35580 AW059918 AA054563 A1280942 T27619 BE621435 N66010 AW589527 A1160414 AA283090 AA962536 H82726 W52115 W45432 W60433 AA577548 AA146714 BE150994 AA054615 AW796025 AW382768 BE565671 C00444
75	100668	26401_3	AA054555 L05424 X56794 S66400 X55150 W60071 AW351820 X55938 M83326 BE005289 BE070059 M83324 BE005248 BE069717 BE181648 BE069700 AW606203 BE069721 AW382138 AW803776 BE463954 BE005334 BE005274 T27386 AA932714 AA972695 AW377728 AI632506 T29066 AI783934 AW377727 BE163715 AL047291 AA279047 AA523003 BE008048 BE440141 W23614 BE090519 BE092193 N29181 N20358 N44153
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00	100491 BE277805	AA147951 AA60	3113 BE253293 AI246588 AI183405 AI954174 AI126891 AI829101 AI123832 AW129670 AA471268 AW170242 AW873079 AA148011 AI608620 AA482961 AI003658 H43261 AA657978 AI735072 R83138 AA722002 AA626271 AW273877 BE464626 AA071483 AA429973 AA494342
			AA620436 AA775597 AA775601 AA826847 A1192585 AA826359 AA411159 A1193419 A1204013 AA705323 AA716255 A1784611 A1081144 A1128227 AA828464 A1148911 A1493446 A1626084 A1189180 A1721196 A1190618 AA284987 A1128543 AA632064 A1333073 A1278470 AA131688
65			A120227 AA020404 A1146311 A1493446 A1020064 A1163160 A1721130 A113010 AA204367 A1120345 AA032004 A1030170 A1710006 A1491768 AA937581 AA630065 AA834257 AW249841 AA583742 A1309756 AA961676 A1760860 AA557818 AA954238 H43655 A1302564
			AA127545 AI609219 H20426 AI042292 AI056466 AA581836 W47002 AA422057 AA937673 F29757 AA829208 AW327462 AA372098 W02144 AA036805 AA487365 AA961037 AI139946 AA487250 AA737118 AI952504 AI242293 AA650552 AI708401 AI633133 AA630848 AA654317 F24128
			A1434165 W46252 AW043879 A1033763 F37228 AA687809 N49087 AA876981 AA506947 A1914572 A1833284 F22253 AA026222 R50166 A1219267 N27095 AA496512 A1784222 A1289904 AA513146 AA528547 AA418700 F36721 A1880700 A1601170 A1862851 A1708633 AA524499
70			AA642220 AA496628 AI718709 W80579 AI720547 F20718 AA649943 AA588229 N40503 H45029 BE262669 BE391069 BE537538 AI510751
			AI906968 AI318611 H46099 AI472604 T60667 AA373087 W32479 AA514034 BE619183 AA134672 AA127544 H26942 BE536689 AW327461 AA422139 AW262357 AW327348 F33510 AI630382 AW827126 F27133 AI335189 AW517599 W80471 AA885814 N89681 BE393173 AA617760
			AA584268 AA460537 AA446261 H20425 N64040 AW276801 AA316367 AA071232 BE545409 AA308292 BE274447 AA380861 AA340038
75			AA341806 AA865579 Al018634 Al766314 Al919302 AA872367 AA991404 Al906961 AA888375 BE621012 AA505388 AA935192 AA290828 R50220 H50814 H44721 AW951723 AA514796 AA418708 AW673377 AA379622 AA977995 AA708224 AA708216 Al318249 Al318233 AA411160
, ,			AA026221 AA316774 AA486908 AI500094 AA096362 AW583742 BE536422 BE618653 R70203 AA131732 AA345048 BE562720 T28342
	100518		NM_004415 AL031058 M77830 BE149760 AW752599 AW848723 AW376697 AW376817 AW376699 AW848371 AW376782 AW848789 AW361413 AW849074 AW997139 AW799304 AW799309 BE077020 BE077017 BE185187 AW997196 BE156621 BE179915 BE006561 BE143155
90			AW899985 BE002107 AW103521 AA857316 AW383133 BE011378 AW170253 BE185750 AW886475 BE160433 J05211 BE082576 BE082584
80			BE004047 AW607238 AW377700 AW377699 BE082526 BE082505 BE082507 BE082514 AW178000 AW177933 AI905935 AW747877 AW748114 BE148516 AW265328 AW847678 AW847688 AW355151 AW365151 AW3
			BE005272 AW365145 BE001925 BE182166 BE144243 BE001923 AI951766 AI434518 BE184920 BE184933 AI284090 BE184941 AW804674 BE184924 C04715 W39488 AW995615 BE184948 BE159646 AW606653 AA099891 AA131128 AA337270 AA340777 AW384371 AA852212
85			R58704 AW366566 AW364859 AA025851 AA025852 AA455100 AA719958 AW352220 AW996245 BE165351 BE073457 AA377127 AW890264
O.J			AW609750 AW391912 AW849590 T87267 AW853812 AA852213 W74149 BE009090 AA056401 H91011 AW368529 AW390272 C18467

	W	02/086443 PCT/US02/12476	
5		AW574920 N57176 AA026480 AW576767 H93284 AA026863 AW177787 AA026654 AW177786 BE092134 BE092137 BE092136 AW177784 Al022862 BE091653 AW376811 AW848592 AA040018 BE185331 BE182164 AA368564 AW951576 T29918 AA131077 W95048 W25458 AW205789 H90899 N29754 W32490 R20904 BE167181 BE167165 N84767 H27408 H30146 A190590 C03378 A155403 A1025263 AA128470 Al392926 AF139065 AW370813 AW370827 AW798417 AW798780 AW798883 AW798569 R33557 AA149190 C03029 AW177783 AA088656 AW370829 AA247685 BE002273 AJ760816 A4339101 AW879451 AJ700963 AA451923 AJ340326 AJ590975 T48793 AJ58096 A142882 AA03997: AI470146 AA946936 BE002273 AJ760816 AJ439101 AW879451 AJ700963 AA451923 AJ340326 AJ590975 T48793 AJ58096 A142882 AA03997: AI470146 AA946936 BE067737 BE067786 W19287 AA644381 AA702424 AJ417612 AJ305554 AJ686869 AJ568892 AW190555 AJ571075 AJ22057: AA056527 AJ471874 AJ304772 AW517828 AJ915596 AJ627383 AJ270345 AW021347 AW166807 AW105614 AJ346078 AA552300 W95070	3
10		A1494059 A1911702 AA149191 AA026864 A1830049 A1887258 AW780435 A1910434 A1819984 A1858282 A1078449 A1025932 A1860584 A1635878 AA026047 AA703232 D12062 AW192085 AA658154 AW514597 AW591892 T87181 AA782066 AW243815 AW150038 AW268383 AW004633 A1927207 AA782109 AW473233 A1804485 AW169216 A1572669 AA602182 AW015480 AW771865 A1270027 AA961816 AA283207 A1076962 A1498487 A1348053 A1783914 H44405 AW799118 AA128330 AA515500 AA918281 W02156 A1905927 AA022701 W38382 R20795 T77861 AW860878	
15	100528	5979_1 BE386801 AU077299 AA143755 BE302747 AA853375 U30162 BE274163 BE277479 BE408180 BE274874 C15000 AA047476 N27099 Al359165 Al638794 Al151283 Al863925 AW444977 Al207392 AA931283 AA443112 R40138 AW068538 AA351008 AA676972 R62503 AA916492 AW00186 H42334 H38280 AA121497 AA114137 Al755038 M17783 AA38786 BE274462 AL753182 C05975 AA347404 AW069298 Al754351 Al754044 AA188808 AA186879 AA565243 AL040655 AA456177 Al750722 AA045756 AA213580 C16936 AW578747 AW753731 H41632 N44761 R58560 R61260 AA039902 N59721 AW992543 R68380 AA149686 T29017 H03739 BE383822 BE387105 BE408251 BE410425 H41560 AA247591 BE389677 Al752233 Al566195 AA868004 Al424523 AW753720 AA852159 BE386803	
20	100559	260_1 NM_00094 L02870 D13694 S51236 M96984 AW946290 M65158 AI285422 D29523 AL119886 AW630655 L06862 Al884355 AW168737 T29085 AW797005 AW801340 AI355504 AW079048 AW801337 AI690455 AI972063 AW268565 W68588 AA587326 AA883498 AI033523 AW510356 AW591998 H98463 AL043852 AI150055 AI566239 AI624803 AA844717 H40670 AA922334 AI864424 AW615094 AW451233 AI302203 F31221 AI872170 W68589 AA904478 AI917631 AW014208 AW450759 AA847625 AI284033 AA848176 AA598507	
25	100576 124357 101624 101625 135158	986_1 X00356 NM_001741 M26095 X03662 M12667 X02330 X02330 AA716058 AW296074 X04861 Al695720 AA719597 enbank_N22401 N22401 ntrez_M55998 M55998 ntrez_M57293 M57293 7963_1 AL037551 Al804716 AW439811 Al569470 AA075299 Al738572 Al270388 Al816783 AW263026 Al633951 Al655285 Al990572 Al950425 AW241533 AA916883 AA576693 AA160156 AA613783 AW078884 Al888282 Al275241 Al133467 AA164921	

Table 2A shows 504 genes down-regulated in lung tumors relative to normal lung and chronically diseased lung. Chronically diseased lung samples represent chronic normalignant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 59660 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression. 5

expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRN/								f mRNA ex	pression.			
	Pkey: ExAccn: Unigeneli	Exempl	Eos probeset ar Accession (a number	ldentifier number number, Genbank accession number								
10	Unigene 1 R1:	Fitle: Unigeno 90th pe	e gene title rcentile of AI f	or normal lung samples divided by the 80th percer								
	R2:	nedian	of Al for norm	al lung samples divided by 90th percentile of Al fo	r adenocarcin	oma and s	quamous	cell carcir	roma lung	tumor sar	nples.	
15	R3:	median	of Al for norm	al lung samples minus the 15th percentile of Al for Al for adenocarcinoma and squamous cell carcin	. 91) uouva ini	ng, cnronic	ally disea	iseo lung a na 15th ner	uno lumor rentile of	Al for all o	ormal	
15		the 90th	percentile of	Al for adenocarcinoma and squamous ceu carcin ased lung and tumor samples.	oma lung tum	or samples	Hillius u	ia ioui bei	Cerillie O	M IOI GII II	Dittion	
	R4:	average	of Al for nor	nal lung samples divided by average Al for squam	ous cell carcir	noma and a	denocan	cinoma lun	g tumors.			
	R5:	modion	of Al for norm	all near complex divided by the QNH nementile of A	u for adenoca	ircinomas.						
20	R6:	median	of Al for norm	al lung samples minus the 15th percentile of Al for	all normal lu	ng, chronic	ally disea	sed lung a	ind tumor	samples d	rvided by t	ne sum
20	07.	percent	ile of Al for ad	enocarcinomas minus the 15th percentile of Al for nal lung samples divided by the 90th percentile of	ali normai iun Al for souamo	ig, chronica vis cell car	any uisea cinomas.	sea lung a	nu tuntor.	sampies.		
	R7: R8:	median	of Al for norm	al lung samples minus the 15th percentile of Al for	r ali normai iui	no. Chronic	aliy disea	isea lung a	and tumor	samples d	ivided by t	he 90th
		percent	ile of A) for sq	uamous cell carcinomas minus the 15th percentile	of Al for all n	ormal lung	, chronica	ally disease	ed lung an	id tumor sa	mples.	*
25	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2	R3	R4	R5	R6	R7	R8
23	,				40.20							
	100095 100115	Z97171 NM_002084	Hs.78454 Hs.336920	myocilin; trabecular meshwork inducible glutathione peroxidase 3 (plasma)	40.20							3.46
	100138	U83508	Hs.2463	angiopoietin 1			2.30					
30	100299	D49493	Hs.2171	growth differentiation factor 10		11.00	•			2.00		
	100305	U86749	Hs.80598	transcription elongation factor A (SII);						3.06		3.16
	100447 100458	NM_014767	Hs.74583 Hs.247979	KIAA0275 gene product Vpre-B	42.40							0.10
	100456	S74019 AA005247	Hs.285754	Hepatocyte Growth Factor Receptor						4.13		`
35	100959	AA359129	Hs.118127	actin: alpha: cardiac muscle				125.60				
	101032		Hs.46039	phosphoglycerate mutase 2 (muscle)	36.40			34.60		•		
	101081 101088	AF047347 X70697	Hs.4880 Hs.553	amyloid beta (A4) precursor protein-bind solute carrier family 6 (neurotransmitte	•	•		193.20				
		AJ250562	Hs.82749	transmembrane 4 superfamily member 2						3.10		
40	101180	U11874	Hs.846	interleukin 8 receptor; beta				54.86				
	101308	L41390	11- 00004	"Homo sapiens core 2 beta-1,6-N-acetylgl	33.20			36.40				
	101330 101345	L43821 NM_005795	Hs.80261 He 152175	enhancer of filamentation 1 (cas-like do Calcitonin receptor-like			2.29	00.40	-			
	101346	Al738616	Hs.77348	hydroxyprostaglandin dehydrogenase 15-(N				70.55				
45	101397	M26380	Hs.180878	lipoprotein lipase							201	3.54
	101414	NM_000066		complement component 8; beta polypeptide	• "			34.60	٠.	٠.	3.81	
	101435 101507	NM_001100 X16896	Hs.1288 Hs.82112	actin; alpha 1; skeletal muscle interleukin 1 receptor; type l				37.60				
	101530	M29874	Hs.1360	cytochrome P450; subfamily IIB (phenobar								4.25
50	101537	Al469059	Hs.184915	zinc finger protein; Y-linked			2.54					
		NM_000102		cytochrome P450; subfamily XVII (steroid	39.40	5.50				´- ·	,	
	101545	BE246154 BE207611	Hs.154210 Hs.123078	EDG1; endothelial differentiation, sphin thyrold stimulating hormone receptor	35.40	13.00						*
	101560		Hs.83733	Intercellular adhesion molecule 2, exon								3.38
55	101574	M34182	Hs.158029	protein kinase; cAMP-dependent; catalyti						4.37		3.80
	101605 101621	M37984 BE391804	Hs.118845 Hs.62661	troponin C; slow guanylate binding protein 1; interferon-	30.20	,			•			0.00
		AA299330	Hs.1042	Sjogren syndrome antigen A1 (52kD; ribon					•		2.75	
	101829	AW452398	Hs.129763	solute carrier family 8 (sodium/calcium				00.00		3.37		
60	101842	M93221	Hs.75182	mannose receptor; C type 1			2.32	38.20				
	101961 101994	AW004056 T92248	Hs.168357 Hs.2240	"Hs-TBX2=T-box gene {T-box region} [huma uteroglobin			2.02					6.85
	102020	AU077315	Hs.154970	transcription factor CP2			2.45					0.75
<i>(</i> =	102091	BE280901	Hs.83155	aldehyde dehydrogenase 7	E4.00							6.75
65	102112	AW025430	Hs.155591	forkhead box F1	54.60							3.98
	102190 102202	AA723157 NM_000507	Hs.73769 Hs.574	folate receptor 1 (adult) fructose-bisphosphatase 1	,					•		3.62
	102241		Hs.268107	Multimerin	·		2.32					
70	102310	U33839		Accession not listed in Genbank	. 00.40	7.00						
70		U41898 U60115	Nº 330000	"Human sodium cotransporter RKST1 mRNA, "Homo sapiens skeletal muscle LIM-protei	29.40							3.75
	102571 102620	AA976427	Hs.239069 Hs.121513	Human clone W2-6 mRNA from chromosome X						3.07		
	102636	U67092		"Human ataxia-telanglectasia locus prote			2.40					
75	102667	U70867	Hs.83974	solute carrier family 21 (prostaglandin			3.15			3.56		
75	102675 102698	U72512 M18667	Hs.7771 Hs.1867	"Human B-cell receptor associated protei propastricsin (pepsinogen C)						0.00		4.51
	102698	W10007 U79251	Hs.99902	opioid-binding protein/cell adhesion mol				•	12.00			
	102852	V00571	Hs.75294	corticotropin releasing hormone	37.40				40.00			
00	103026	X54162	Hs.79386	thyrold and eye muscle autoantigen D1 (6	90 00				13.00			
80	103028 103098	X54380 M86361	Hs.74094	pregnancy-zone protein Human mRNA for T cell receptor; clone IG	28.80				10.00			
	103098		Hs.295449	parvalbumin		6.00						
	103241	X76223		H.sapiens MAL gene exon 4			2.47					
	103280	U84722	Hs.76206	Cadherin 5, VE-cadherin (vascular epithe			2.69					

H.sapiens MAL gene exon 4 Cadherin 5, VE-cadherin (vascular epithe

keratin; hair, acidic; 5

2.16

103280

103360

85

U84722

Y16791

Hs.76206

Hs.73082

	w	O 02/08	6443							PCT/	US02/:	12476
	103496	Y09267	Hs.132821	flavin containing monooxygenase 2								5.97
	103508	Y10141	Un 4.49494	*H.sapiens DAT1 gene, partial, VNTR*			2.40			3.27		
	103561 103569	NM_001843 NM_005512		contactin 1 glycoprotein A repetitions predominant			2.99					
5	103575	Z26256		"H.sapiens isoform 1 gene for L-type cal						4.18		
	103627	Z48513 BE244667	Un 2021EE	H.sapiens XG mRNA (clone PEP6) CGI-100 protein						3.44	2.25	
	103767 103850	AA187101	Hs.296155 Hs.213194	Hypothetical protein MGC10895; sim to SR				46.55				
10	104078	AA402801	Hs.303276	ESTs						3.05		
10	104326	AW732858	Hs.143067	ESTs dual-specificity tyrosine-(Y)-phosphoryl						3.54 3.16		
	104352 104398	BE219898 Al423930	Hs.173135 Hs.36790	ESTs; Weakly similar to putative p150 (H	64.80					•		
	104473	A1904823	Hs.31297	ESTs			0.43					3.38
15	104493	AW960427	Hs.79059 Hs.292979	ESTs; Moderately similar to TGF-BETA REC ESTs	28.60		2.47					
13	104495 104595	AW975687 Al799603	Hs.271568	ESTs	20.00					3.42		
	104597	A1364504	Hs.93967	ESTs; Weakly similar to Slit-1 protein [6.00	•					
	104659	AW969769	Hs.105201	ESTs ESTs	34.00	11.00						
20	104686 104691	AA010539 U29690	Hs.18912 Hs.37744	ESTs; Beta-1-adrenergic receptor	56.80	11.00						
	104764	A1039243	Hs.278585	ESTs			·	60.40				
	104776	AA026349 AA035613	Un 4.44000	ESTs	34.20		3.03					
	104825 104865	T79340	Hs.141883 Hs.22575	Homo sapiens cDNA: FLJ21042 fls, clone C	41.20		0.00					
25	.104942	NM_016348	Hs.10235	ESTs				40.00				3.27
	104989	R65998	Hs.285243	ESTs				40.00				3.20
	105062 105101	AW954355 H63202	Hs.36529 Hs.38163	ESTs	34.20							
20	105173	U54617	Hs.8364	ESTs		40.00						4.17
30	105194	R06780	Hs.19800	ESTs ESTs		16.00	2.34					
	105226 105256	R58958 AA430650	Hs.26608 Hs.16529	transmembrane 4 superfamily member (tetr			2.72					
	105394	BE245812	Hs.8941	ESTs	00.00		2.61					
35	105647	Y09306	Hs.30148	homeodomain-interacting protein kinase 3	33.60							3.59
33	105789 105817	AF106941 AA397825	Hs.18142	arrestin; beta 2 synaptopodin						4.46	·	5.55
	105847	AW964490	Hs.32241	ESTs			0.40	35.40			٠.	
	105894	A1904740	Hs.25691	calcitonin receptor-like receptor activi		7.00	3.43					
40	105999 106075	BE268786 AA045290	Hs.21543 Hs.25930	ESTs ESTs		7.00		42.60				
	106178	AL049935	Hs.301763	KIAA0554 protein	34.80				40.00			
	106381	AB040916	Hs.24106 Hs.154162	ESTs ADP-ribosylation factor-like 2					12.00	3.69		
	106467 106536	AA450040 AA329648	Hs.23804	ESTs				96.40				
45	106569	R20909	Hs.300741	sordin				47.20				
	106605 106842	AW772298 AF124251	Hs.21103 Hs.26054	Homo sapiens mRNA; cDNA DKFZp564B076 (fr novel SH2-containing protein 3			2.55	220.40				
	106844		Hs.158213	sperm associated antigen 6	39.20							
50	106870	AI983730	Hs.26530	serum deprivation response (phosphatidy)			2.28					4.28
50	106943 106954	AW888222 AF128847	Hs.9973 Hs.204038	ESTs ESTs								4.32
era amen era jaga algere	107106	AA862496	Hs.28482	ESTs					10.45	* :		
	107163	AF233588	Hs.27018	ESTs			2.57	•		3.84		
55	107201 107238	D20378 D59362	Hs.30731 Hs.330777	EST EST		8.00				3.04	•	
55	107376	U90545	Hs.327179	solute carrier family 17 (sodium phospha		10.67						
	107530	Y13622	Hs.85087	latent transforming growth factor beta b			2.32	34.60				
	107688 107706	AW082221 AA015579	Hs.60536 Hs.29276	ESTs ESTs	28.40			34.00				
60	107723	AA015967	1101-0210	EST						3.29		
	107727	AA149707	Hs.173091	DKFZP434K151 protein				80.80 51.40				
	107750 107751	AA017291 AA017301	Hs.60781 Hs.235390	ESTs ESTs	•			J 11.TV		3.14		
	107873	AK000520	Hs.143811	ESTs		9.00				0.05		
65	107899	BE019261 AA036811	Hs.83869 Hs.48469	ESTs; Wealdy similar to !!!! ALU SUBFAMI ESTs				44.60		3.65		
	107994 107997	AL049176	Hs.82223	Human DNA sequence from clone 141H5 on c				32.00				
	108041	AW204712	Hs.61957	ESTs				30.80			4.75	
70	108048	AI797341 AA070773	Hs.165195	ESTs "zm53g11.s1 Stratagene fibroblast (#9372			2.33				4.75	
70	108338 108434	AA078899		"zm94b1.s1 Stratagene colon HT29 (#93722			2.00				2.92	
	108447	AA079126		"zm92a11.s1 Stratagene ovarian cancer (#				24.00		3.06		
	108480 108499	AL133092 AA083103	Hs.68055	ESTs *zn1b12.s1 Stratagene hNT neuron (#93723				34.00				3.36
75	108535	R13949	Hs.226440	Homo sapiens clone 24881 mRNA sequence					19.00			
	108550	AA084867		"zn11f6.s1 Stratagene hNT neuron (#93723			2 22		12.00			
	108604 108625	AA934589 AW972330	Hs.49696 Hs.283022	ESTs ESTs			2.33					5.82
0.0	108629	AA102425	·	"zn24c6.s1 Stratagene neuroepithelium NT							3.42	
80	108655	AA099960	Un 44=00=	"zm65c6.s1 Stratagene fibroblast (#93721		7.00 6.05						
	108756 108864	AA127221 Al733852	Hs.117037 Hs.199957	Homo sapiens mRNA; cDNA DKFZp564N1164 (f ESTs	28.80	0.03						
	108895	AL138272	Hs.62713	ESTs	32.80							
85	108921	A1568801	Hs.71721	EST8	28.80			57.80				
65	108967	AA142989	Hs.71730	ESTs	20.00							

	W	O 02/086	6443							PCT/	U S02/ :	12476
	109001	AJ056548	Hs.72116	ESTs, Moderately similar to hedgehog-int			2.57				2.11	
	109003 109004	AA147497 AA156235	Hs.71825 Hs.139077	ESTS EST		5.60					2.11	
	109065	AA161125	Hs.252739	EST					10.00			
5	109250	H83784	Hs.62113	ESTs; Weakly similar to PHOSPHATIDYLETHA							3.44 2.92	
	109490 109510	AA233416 Al798863	Hs.139202 Hs.87191	ESTs ESTs			2.40				2.32	
	109578	F02208	Hs.27214	ESTs		10.00						
10	109601	F02695	Hs.311662	EST				40.80 54.40				
10	109613 109550	H47315 R31770	Hs.27519 Hs.23540	ESTs ESTs	31.20			54.40		•		
	109682	H18017	Hs.22869	ESTs	0.1.20	8.40						
	109724	D59899	Hs.127842	ESTs				29.40	8.00			
15	109782 109833	AB020644 R79864	Hs.14945 Hs.29889	long fatty acyl-CoA synthetase 2 gene ESTs		10.00			0.00			
13	109837	H00656	Hs.29792	ESTs			6.49					
	109977	T64183	Hs.282982	ESTs				407.00			2.75	
	109984 110146	Al796320 H41324	Hs.10299 Hs.31581	ESTs ESTs; Moderately similar to SYNTAXIN 1B				107.00			2.22	
20	110271	H28985	Hs.31330	ESTs						3.48		
	110280	AW874263	Hs.32468	ESTs	44.20			00.00				
	110420	R93141	Hs.184261 Hs.11038	ESTs ESTs	28.40			32.00				
	110578 110634	T62507 R98905	Hs.35992	ESTs .	20.40				20.00		•	
25	110726	AW961818	Hs.24379	potassium voltage-gated channel; shaker-								4.15
	110837	H03109	Hs.108920	ESTs; Weakly similar to semaphorin F [H.			3.13	56.80				
	110875 110894	N35070 R92356	Hs.26401 Hs.66881	tumor necrosis factor (ligand) superfami ESTs; Moderately similar to cytoplasmic		5.33	3.10					
	110971	AI760098	Hs.21411	ESTs				44.60				
30	111023	AV655386	Hs.7645	ESTs FOT	32.40				17.14			
	111057 111247	T79639 AW058350	Hs.14629 Hs.16762	ESTs Homo sapiens mRNA; cDNA DKFZp564B2062 (f					(7.14		4.58	
	111330	BE247767	Hs.18166	KIAA0870 protein								3.42
25	111374	BE250726	Hs.283724	ESTs; Moderately similar to HYA22 [H.sap				33.20				3.91
35	111442 111737	AW449573 H04607	Hs.181003 Hs.9218	ESTs ESTs				53.20				
	111747	Al741471	Hs.23666	ESTs	46.20							
	111807	R33508	Hs.18827	ESTs		16.00			•	3.91		
40	111862 112045	R37472 Al372588	Hs.21559 Hs.8022	EST TU3A protein						3.91	2.74	
40	112043	R43713	Hs.22945	EST							4.92	
	112214	AW148652	Hs.167398	ESTs			0.40		13.00	. 3		
	112263 112314	R52393 AW206093	Hs.25917 Hs.748	ESTs ESTs		9.00	2.43					
45 ·	112324	R55965	Hs.26479	limbic system-associated membrane protei		0.00			14.00			
	112362	AW300887	Hs.26638	ESTs; Weakly similar to CD20 receptor [H			2.49					
	112380 112425	H63010 AA324998	Hs.5740 Hs.321677	ESTs ESTs: Weakly similar to !!!! ALU SUBFAMI		8.00	2.34					
	112473	R65993	Hs.279798	pregnancy specific beta-1-glycoprotein 9		0.00				4.53		
50 .	112492	N51620	Hs.28694	ESTs				29.80		2 62		
	112541 112620	AF038392 R80552	Hs.116674 Hs.29040	ESTs ESTs		•	2.37			3.62		
	112623	AW373104	Hs.25094	ESTs			2.26					
<i>E E</i>	112867	T03254	Hs.167393	ESTs .		6.50			12.00			
55	112894 112954	T08188 AA928953	Hs.3770 Hs.6655	ESTs ESTs		6.50 7.00						
	113029	AW081710	Hs.7369	ESTs; Weakly similar to IIII ALU SUBFAMI								4.39
	-113086	AA346839	Hs.209100	DKFZP434C171 protein				•	40.00			4.47
60	113140 113252	T50405 NM_004469	Hs.175967 Hs.11392	ESTs c-fos induced growth factor (vascular en	-	14.00			10.00			
. 00	113257	Al821378	Hs.159367	ESTs						3.72		
	113394	T81473	Hs.177894	ESTs	25.00					3.60		
	113437 113454	T85349 Al022166	Hs.15923 Hs.16188	EST ESTs	35.00	6.00						
65	113502	T89130	113.10100	ESTs	39.60	0.00						
	113552	Al654223	Hs.16026	ESTs							2.58	3.88
	113645 113691	T95358 T96935	Hs.333181 Hs.17932	ESTs EST				38.20			2.30	
	113706	AA004693	Hs.269192	ESTs						3.09		
70	113883	U89281	Hs.11958	oxidative 3 alpha hydroxysteroid dehydro	20.40		2.31					
	113924 114035	BE178285 W92798	Hs.170056 Hs.269181	Homo sapiens mRNA; cDNA DKFZp586B0220 (f ESTs	30.40				13.00		-	
	114058	AK002016	Hs.114727	ESTs								5.00
75	114084	AA708035	Hs.12248	ESTs			2 24	40.60				
75	114121 114124	H05785 W57554	Hs.25425 Hs.125019	ESTs Human lymphoid nuclear protein (LAF-4)		7.00	2.31					
	114275	AW515443	Hs.306117	interleukin 13 receptor; alpha 1		6.00						
	114297	AA149707	Hs.173091	DKFZP434K151 protein				48.80		3.45		
80	114427 114449	AA017176 AA020736	Hs.33532	ESTs; Highly similar to Miz-1 protein (H "ze63b11.s1 Soares retina N2b4HR Homo sa					10.00	3.43		•
	114452	Al369275	Hs.243010	ESTs, Moderately similar to RTC0_HUMAN G		14.00						
	114609	AA079505		"zm97a5.s1 Stratagene colon HT29 (#93722				35.40		3.13		
	114648 114731	AA101056 BE094291	Hs.155651	"zn25b3.s1 Stratagene neuroepithelium NT Homo sapiens HNF-3beta mRNA for hepatocy				JJ.40		-		3.42
85	114762	AA146979	Hs.288464	ESTs	33.00							

	w	O 02/086	5443							PCT/	US02/1	2476
	114776	AA151719	Hs.95834	EST ₈	34.40							
	115009	AA251561	Hs.48689	EST8	30.20							
	115272 115279	AW015947 AW964897	Hs.290825	ESTs; Weakly similar to hypothetical L1 ESTs	32.60	6.00						
5	115302	AL109719	Hs.47578	ESTs		0.00			12.00			
•	115365	AW976252	Hs.268391	ESTs						3.32		
	115559	AL079707	Hs.207443	ESTs				48.00				
	115566	AI142336	Hs.43977	ESTs ESTs, Weakly similar to (defline not ava	31.40			56.20				
10	115683 115744	AF255910 AA418538	Hs.54650 Hs.43945	ESTs; Highly similar to dJ1178H5.3 [H.sa	31.40			33.60				
	115819	AA486620	Hs.41135	Endomucin 2				74.40				
	115949	A1478427	Hs.43125	ESTs			3.18	200.00				
	115965	AA001732	Hs.173233	ESTs ESTs				388.80 33.20				
15	116035 116049	AA621405 AA454033	Hs.184664 Hs.41644	ESTs				45.80				
-0	116081	AI190071	Hs.55278	ESTs						3.57		
	116082	AB029496	Hs.59729	ESTs	C0.00		3.06					
	116213	AA292105	Hs.326740 Hs.50841	leucine rich repeat (in FLII) interactin ESTs; Weakly similar to tuftelin [M.musc	50.60		3.85					
20	116228 116250	A1767947 N76712	Hs.44829	ESTs		6.00	0.00					
	116419	Al613480	Hs.47152	ESTs; Weakly similar to testicular tekli		_		30.00				
	116617	D80761	Hs.45220	EST	47.00	•	2.27					
•	116784	AB007979	Hs.301281 Hs.38218	tenascin R (restrictin; janusin) ESTs	47.20			41.20				
25	116835 116970	N39230 AB023179	Hs.9059	KIAA0962 protein					11.00			•
	117023	AW070211	Hs.102415	ESTs				91.00		•		
	117027	AW085208	Hs.130093	ESTs	49.40			22.60				
	117036	H88908	Hs.41192	EST ESTs		8.67		32.60				
30	117110 117209	AA160079 W03011	Hs.172932 Hs.306881	ESTs		0.01		30.60				
	117325	N23599	Hs.43396	ESTs					9.29			
	117454	N29569	Hs.44055	ESTs	44.00				•	3.19		
	117475	N30205 BE219453	Hs.93740	ESTS	44.00	16.00						
35	117543 117567	AW444761	Hs.42722 Hs.44565	ESTs ESTs		10.00			12.00			
70	117570	N48649	Hs.44583	ESTs					11.00		•	
	117600	N34963	Hs.44676	EST		c 00				3.74		
	117730	N45513 N48325	Hs.46608 Hs.93956	ESTs EST		6.00 9.00						
40	117791 117929	N51075	Hs.47191	ESTs		0.00		29.20				
	117990	AA446167	Hs.47385	ESTs		8.00						
	118224	N62275	Hs.48503	EST	31.40							
	118244 118357	N62516 AL109667	Hs.48556 Hs.124154	ESTs Homo sapiens mRNA full length insert cDN	32.80		2.40			•		
45	118446	N66361	Hs.269121	ESTs	•		2.28					
	118447	N66399	Hs.49193	EST	30.80							
	118530	N67900	Hs.118446	ESTs						3.10 3.41		
	118549 118823	N68163 W03754	Hs.322954 Hs.50813	EST ESTs; Weakly similar to long chain fatty			3.94			0.41		
50	118862	W17065	Hs.54522	ESTs						3.58		
	118935	Al979247	Hs.247043	KIAA0525 protein				33.00	44.40			·
	118944	AI734233	Hs.226142	ESTs; Weakly similar to !!!! ALU SUBFAMI		14.00			11.43			
	118995 119073	N94591 BE245360	Hs.323056 Hs.279477	ESTs ERG-2/ERG-1; V-ets avian erythroblastosi		14.00		52.60				
55	119268	T16335	Hs.65325	EST	31.40							•
	119514	W37937		Accession not listed in Genbank			0.75			3.50		
	119824	W74536	Hs.184 Hs.58419	advanced glycosylation end product-speci - DKFZP586L2024 protein			2.75					3.21
	119831 119861	AL117664 W78816	Hs.49943	ESTs; Moderately similar to !!!! ALU SUB				33.80				••
60 -	119889	W84346	Hs.58671	ESTs				30.03				
	119921	W86192	Hs.58815	ESTs	29.00					3.80		
	120082 120094	H80286 AA811339	Hs.40111 Hs.124049	ESTs .		6.00				5.00		
	120034	W57554	Hs.125019	Human lymphoid nuclear protein (LAF-4)		5.00		36.60	-			
65	120378	AA223249	Hs.285728	ESTs		12.00						
	120404	AB023230-	Hs.96427	KIAA1013 protein	39.40				8.00			
	120504 120512	AA256837 N55761	Hs.194718	ESTs ESTs	33.00				0.00			
	120667	AA287740	Hs.78335	microtubule-associated protein; RP/EB fa								4.18
70	120777	AA287702	Hs.10031	KIAA0955 protein				46.60				
	121082 121191	AA398722 AA400205	Hs.104447	ESTs ESTs	41.60			39.00				
	121248	AA400914	Hs.97827	EST .	11.00						5.08	
~-	121363	A1287280	Hs.97933	ESTs					12.00			
75	121366	AI743515	11- 00	ESTs					20.00	3.32		
	121483 121518	Al660332 AA412155	Hs.25274	ESTs; Moderately similar to putative sev ESTs				30.20		U.UZ		
	121518	AA412133 AA412442	Hs.98132	ESTS			2.29	J4.20				
0.0	121622	AA416931	Hs.126065	ESTs		9.00						
80	121665	AA416556	Hs.98234	ESTS	24.00			34.80				
	121709 121730	A1338247 A1140683	Hs.98314 Hs.98328	Homo sapiens mRNA; cDNA DKFZp586L0120 (f ESTs	34.80 38.80							
	121740	AA421138	Hs.98334	EST	00.00	7.00						
0.5	121772	Al590770	Hs.110347	Homo sapiens mRNA for alpha integrin bin	36.20							3.61
85	121821	AL040235	Hs.3346	ESTs								J.U I

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	121835	AB033030	Hs.300670	ESTs			2.34					
	121841	AA427794	Hs.104864	ESTs			2.61				2.25	
	121885 121888	AA934883 AA426429	Hs.98467 Hs.98463	ESTs ESTs							2.92	
5	121938	AA428659	Hs.98610	ESTs				46.80				
	121950	AA429515		EST				31.40				
	122030	AA431310	Hs.98724	ESTs	34.40						2 50	
	122054	AA431725	Hs.98746	EST	40.40						3.58	
10	122211 122233	AA300900 AA436455	Hs.98849 Hs.98872	ESTs; Moderately similar to bithoraxoid- EST	49.40 29.80							
10	122247	AA436433 AA436676	Hs.98890	EST	20.00			39.80				
	122253	AA436703	Hs.104936	ESTs; Weakly similar to hypothetical pro		9.00						
	122266	AA435840	Hs.98907	EST						3.60		
1.5	122285	AA436981	Hs.121602	EST	20.00					3.14		
15	122409 122485	AA446830 AA524547	Hs.99081 Hs.160318	ESTs phospholemman	30.80		2.65					
	122697	AA420683	Hs.98321	Homo sapiens cDNA FLJ14103 fis, clone MA		15.00						
	122772	AW117452	Hs.99489	ESTs		6.67						
20	122831	AI857570	Hs.5120	ESTs				00.00		3.37		
20	122913	A1638774	Hs.105328	ESTs				32.20 41.80				
	123049 123076	BE047680 Al345569	Hs.211869 Hs.190046	ESTs ESTs	35.80			41.00				
	123136	AW451999	Hs.194024	ESTs	***************************************		•				2.58	
	123309	N52937	Hs.102679	ESTs					19.00			
25	123455	AA353113	Hs.112497	ESTs				82.80		2.05		
	123691	AA609579	Hs.112724	ESTs	35.40					3.95		
	123756 123802	AA609971 AA620448	Hs.112795	EST Homo sapiens clone 24760 mRNA sequence	58.00							
	123837	AI807243	Hs.112893	ESTs	00.00			32.40				
30	123844	AA938905	Hs.120017	olfactory receptor; family 7; subfamily			2.63					
	123936	NM_004673		ESTs	29.00			70.00				
	123987	C21171	Hs.95497	ESTs; Weakly similar to GLUCOSE TRANSPOR	28.40			70.60				
	124013 124160	AI521936 R40290	Hs.107149 Hs.124685	ESTs; Weakly similar to PTB-ASSOCIATED S ESTs	20.40				13.00			
35	124205	H77570	Hs.108135	ESTs						4.74		•
	124226	AA618527	Hs.190266	ESTs			2.35					
	124246	H67680	Hs.270962	ESTs .		47.00		29.40				
	124348	AI796320	Hs.10299	ESTS		17.00	3.07					
40	124358 124409	AW070211 Al814166	Hs.102415 Hs.107197	"yw35g11.s1 Morton Fetal Cochlea Homo sa ESTs			3.07			3.14		
40	124442	AW663632	Hs.285625	TATA box binding protein (TBP)-associate			2.48			••••		
	124468	N51413	Hs.109284	ESTs				30.80				
•	124479	AB011130	Hs.127436	calcium channel; voltage-dependent; alph								6.03
45	124519	Al670056	Hs.137274	ESTs; Weakly similar to SPLICEOSOME ASSO	59.20		2.50					
43	124711 124866	NM_004657 AJ768289	Hs.304389	serum deprivation response (phosphatidyl ESTs	09.ZŲ	8.00						
	124874	BE550182	Hs.127826	ESTs		0.00		37.60				
	125097	AW576389	Hs.335774	ESTs					10.00			
50	125179	AW206468	Hs.103118	ESTs						3.12	2.79	
50	125200	AW836591	Hs.103156 Hs.102720	ESTs				34.20			2.13	
	125299 125400	T32982 AL110151	Hs.102720	ESTs DKFZP586D0824 protein	29.00			UT.20				
	125810	H00083		aryl hydrocarbon receptor-interacting pr	32.20							
~ ~	126176	BE242256	Hs.2441	KIAA0022 gene product		12.00					• •	
55	126303	D78841		HUM525A05B Human placenta polyA+ (TFuji	25.00			33.60				
	126403	AW629054	Hs.125976 Hs.23964	ESTs; Weakly similar to metalloprotease/ ESTs; Weakly similar to HC1 ORF [M.muscu	35.80			29.80				
	126507 126773	AL040137 AA648284	Hs.187584	ESTs	39.60			20.00				
	127307	AW962712	Hs.126712	ESTs; Weakly similar to plL2 hypothetica	28.80							
60	127462	AA760776	Hs.293977	aa59b04.s1 NCI_CGAP_GCB1 Homo saplens c				34.40				
	127486	AW002846	Hs.105468	ESTs		9.00	2.36					
	127572 127609	AA594027 X80031	Hs.191788 Hs.530	ESTs ESTs			2.30	29,40				
	127832	AW976035	Hs.292396	EST8				37.20				
65	127898	AA774725	Hs.128970	ESTs							4.42	
	128073	AW340720	Hs.125983	ESTs				38.40				
	128101	AA905730	Hs.128254	ESTs		7.33					2.58	
	128149 128212	· NM_012214 W27411	Hs.336920	mannosyl (alpha-1;3-)-glycoprotein beta- glutathione peroxidase 3 (plasma)			3.09				2.30	
70	128333	W68800	Hs.12126	ESTs; Weakly similar to LR8 [H.sapiens]			0.00	34.40				
, ,	128364	N76462	Hs.269152	ESTs; Weakly similar to ZINC FINGER PROT		10.00						
	128426	Al265784	Hs.145197	ESTs							4.31	
	128598	AA305407	Hs.102308	potassium inwardly-rectifying channel; s	31.20			41.60				
75	128634	AA464918 AW271273	He 22767	ESTs; Moderately similar to IIII ALU SUB				41.00 87.00				
, ,	128687 128726	AW271273 Al311238	Hs.23767 Hs.104476	ESTs ESTs				J7.00				4.02
	128773	NM_004131		granzyme B (granzyme 2; cytotoxic T-lymp					9.00			
	128833	W26667	Hs.184581	ESTs								3.76
QΛ	128870	H39537	Hs.75309	eukaryotic translation elongation factor			2.66			3.10		
80	128878 128885	R25513 AF134803	Hs.10683 Hs.180141	ESTs cofilin 2 (muscle)					11.00	3.10		
	128998	W04245	Hs.107761	ESTs; Weakly similar to PUTATIVE RHO/RAC							3.21	
	129000	AA744902	Hs.107767	ESTs; Moderately similar to CaM-KII inhi								3.68
05	129038	AW156903	Hs.108124	ribosomal protein L41	24.00					3.17		
85	129098	AW580945	Hs.330466	ESTs	34.60		•					

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	129210	AL039940	Hs.202949	KIAA1102 protein								4.09
	129240 129262	AA361258 BE222198	Hs.237868 Hs.109843	interleukin 7 receptor ESTs			2.29			3.30		
_	129301	AF182277	Hs.330780	Human cytochrome P450-IIB (hIIB3) mRNA;						•		4.05
5	129331 129381	AW167668 AW245805	Hs.279772 Hs.110903	ESTs; Highly similar to CGI-38 protein [claudin 5 (transmembrane protein defeted			2.93					4.09
	129565	ΧΤΤΤΤΤ	Hs.198726	vasoactive intestinal peptide receptor 1			2.50	160.80				
	129595	U09550	Hs.1154	oviductal glycoprotein 1; 120kD ESTs; Weakly similar to collagen alpha 1					10.00	3.40		
10	129613 129782	AW978517 AW016932	Hs.172847 Hs.104105	EST Strong similar to contagen alpha i		9.00				3.40		
	129950	F07783	Hs.1369	decay accelerating factor for complement				87.80				
	129958 129959	R27496 AL036554	Hs.1378 Hs.274463	annexin A3 defensin; alpha 1; myeloid-related seque			2.72	44.60				
1.5	130160	AA305688	Hs.267695	UDP-Gal:betaGlcNAc beta 1;3-galactosyltr				42.20				
15	130259 130273	NM_000328 AW972422	Hs.153614 Hs.153863	retinitis pigmentosa GTPase regulator MAD (mothers against decapentaplegic; Dr			2.54	51.60				
	130312	AF056195	Hs.15430	DKFZP586G1219 protein				•		3.16		
	130436 130523	NM_001928 AA999702	Hs.155597 Hs.214507	D component of complement (adipsin) ESTs						4.77		4.11
20	130799	AB028945	Hs.12696	ESTs		6.00						
	130885	NM_005883 AL050295		adenomatous polyposis coli like KIAA0758 protein						3.54		3.50
	131002 131012	AL030293 AL039940	Hs.22039 Hs.202949	KIAA1102 protein		20.00						0.00
25	131031	NM_001650		aquaporin 4	41.20			31.40				
23	131061 131066	N64328 AW169287	Hs.268744 Hs.22588	ESTs; Moderately similar to KIAA0273 [H. ESTs				29.60				
	131082	AI091121	Hs.246218	ESTs; Weakly similar to zinc finger prot					9.00			2.00
	131087 131161	AF147709 AF033382	Hs.22824 Hs.23735	ESTs; Weakly similar to p160 myb-binding potassium voltage-gated channel; subfami						3.14		3.86
30	131179	AA171388	Hs.184482	DKFZP586D0624 protein						3.80		
	131182 131205	AI824144 NM_003102	Hs.23912	ESTs superoxide dismutase 3; extracellular			2.98					3.67
	131277	AA131466	Hs.23767	ESTs			3.15					
35	131281	AA251716	Hs.25227	ESTs				32.20				3.44
33	131282 131285	X03350 Al567943	Hs.4 Hs.25274	alcohol dehydrogenase 3 (class I); gamma ESTs; Moderately similar to putative sev						6.40		J. 1 1
	131355	R52804	Hs.25956	DKFZP564D206 protein		8.00	•					
	131391 131461	AW085781 AA992841	Hs.26270 Hs.27263	ESTs butyrate response factor 2 (EGF-response	28.80	10.00						
40	131487	F13036	Hs.27373	Homo sapiens mRNA; cDNA DKFZp564O1763 (f	20.00						4.03	
	131517 131545	AB037789 AL137432	Hs.263395 Hs.28564	ESTs; Highly similar to semaphorin VIa [ESTs	39.00				11.00			
	131583	AK000383	Hs.323092	ESTs; Weakly similar to dual specificity		-			10.00			
45	131647 131675	AA359615 H15205	Hs.30089 Hs.30509	ESTs ESTs			2.47			3.06		
15	131676	Al126821	Hs.30514	ESTs	45.80					•		
	131708 131717	S60415 X94630	Hs.30941 Hs.3107	calcium channel; voltage-dependent; beta CD97 antigen			2.28					3.78
~~	131756	AA443966	Hs.31595	ESTs ESTS				40.60	9 A			-
50	131762 131821	AA744902 AA017247	Hs.107767 Hs.164577	ESTs; Moderately similar to CaM-KII inhi ESTs			2.87	·				3.67
	131839	AB014533	Hs.33010	KIAA0633 protein	15		2.01				3.48	
	131861	AL096858 Al418006	Hs.184245	KIAA0929 protein Msx2 interacting nuclea ESTs	54.00	•		49.20			,	
55	132015 132070	BE622641	Hs.3731 Hs.38489	ESTs .				34.80				
	132242	AA332697	Hs.42721	ESTs			2.68 4.66					
	132334 132476	AW080704 AL119844	Hs.45033 Hs.49476	lacrimal proline rich protein Homo sapiens clone TUA8 Cri-du-chat regi	34.20							
60		NM_001290		LIM binding domain 2		13.00	2.66					
00	132533 132598	Al922988 X80031	Hs.172510 Hs.530	ESTs collagen; type IV; alpha 3 (Goodpasture		13.00		30.60				
	132619	H28855	Hs.53447	ESTs; Moderately similar to kinesin ligh						4.02 3.18		
	132652 132726	N41739 N52298	Hs.61260 Hs.55608	ESTs ESTs; Weakly similar to cDNA EST yk484g1					11.43	3.10		
65	133028	R51604	Hs.300842	ESTS			2.37				•	
	133071 133120	BE384932 NM_003278	Hs.64313 Hs.65424	ESTs -tetranectin (plasminogen-binding protein			2.27 2.63				•	
	133129	AA428580	Hs.65551	ESTs								5.49
70	133147 133151	AA026533 NM_014051	Hs.66 Hs.94896	Interleukin 1 receptor-like 1 ESTs			6.20			3.69		
,,,	133213	AA903424	Hs.6786	ESTs				31.40		•••		
	133276 133377	AW978439 AJ131245	Hs.69504 Hs.7239	ESTs SEC24 (S. cerevisiae) related gene famil	41.20				9.00			
7.6	133407	AF017987	Hs.7306	secreted frizzled-related protein 1	50.20							
75	133535	AL134030	Hs.284180	protocadherin 2 (cadherin-like 2)						3.72		3.35
	133537 133656	U41518 BE149455	Hs.74602 Hs.75415	aquaporin 1 (channel-forming integral pr Accession not listed in Genbank			2.65					U.UU
	133689	NM_001872	Hs.75572	carboxypeptidase B2 (plasma)				90.80		3.05		
80	133779 133978	T58486 AF035718	Hs.222566 Hs.78061	ESTs transcription factor 21			2.92			J.UJ		
- =	133985	L34657	Hs.78146	platelet/endothelial cell adhesion molec								3.45
	134000 134111	AW175787 Al372588	Hs.334841 Hs.8022	selenium binding protein 1 TU3A protein			4.49					4.05
9.5	134185	AA285136	Hs.301914	Homo saplens mRNA; cDNA DKFZp586K1220 (f				40.00			3.27	
85	134204	AI873257	Hs.7994	ESTs; Weakly similar to CGI-69 protein (40.80				

	W	O 02/086	443							S02/12476
	134641	Al092634	Hs.156114	protein tyrosine phosphatase; non-recept					3.76	
	134677	AA251363	Hs.177711	ESTs				32.20		
	134745	NM_000685		angiotensin receptor 1B		15.00				
_	134749	T28499	Hs.89485	carbonic anhydrase IV			3.05			
)	134786	T29618	Hs.89640	anglopoietin 1 receptor, TEK tyrosine ki				57.80		
	134825	U33749	Hs.197764	thyroid transcription factor 1						3.73
	134978	AI829008	Hs.333383	ficolin (collagen/librinogen domain-cont			2.52			
	135010	N50465	Hs.92927	ESTs				31.60		
10	135053	AW796190	Hs.93678	ESTs					3.21	
10	135081	AF069517	Hs.173993	RNA binding motif protein 6	28.80					2.2
	135091	AA493650	Hs.94367	ESTs						4.24
	135135	AA775910	Hs.95011	syntrophin; beta 1 (dystrophin-associate		8.00				
	135203	C15737	Hs.269386	ESTs					4.31	
4.5	135236	AI636208	Hs.96901	ESTs	43.00					
15	135266	R41179	Hs.97393	Human mRNA for KIAA0328 gene; partial cd					,	6.42
	135346	NM_000928	Hs.992	phospholipase A2; group IB (pancreas)			3.82			
	135378	AW961818	Hs.24379	potassium voltage-gated channel; shaker-			4.15			
	135387	NM_001972	Hs.99863	elastase 2; neutrophil	37.20					
••	135388	W27965	Hs.99865	EST	38.80					
20	135402	L12398	Hs.99922	dopamine receptor D4					4.21	

TABLE 2B shows the accession numbers for those primekeys lacking unigenelD's for Table 2A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

	Pkey:	Unique Eos probeset identifier number nber: Gene cluster number	
30	Accession		
	Pkey	CAT number Accessions	•
0.5	108447	434527 AA079126	
35	108550	120073_1 AA084867 AA084996	
	108655	127522_1 AA099960 AA113013	
	102397	443711 U41898	
	126303	1525933_1 D78841 D78880	
40	125810	1554054_1 H00083 R81062	
40	103627	2615_2 Z48513 Z48512	
	121366	280401_1 AI743515 AA405617 AW276706	•
	114609	116777_1 AA079505 AA079537	
	115272	172113_1 AW015947 AA211890 AA279425	
45	108338 108434	112186_1 AA070773 AA070774 114012_1 AA078899 AA078782 AA075788	
43	123802	genbank_AA620448 AA620448	
	102310	NOT_FOUND_entrez_U33839 U33839	
	102636	entrez_U67092_U67092	
	104776	genbank AA026349 AA026349	
50	120504	genbank_AA256B37 AA256B37	
-	113502	genbank_T89130T89130	
	108499	genbank_AA083103 AA083103	
	10130B	entrez L41390 L41390	
	108629	genbank_AA102425 AA102425	
55	103098	221_215 M86361 Z26593 X02850 D13070 AE000659 M17649 M87869	M87871 X61077 M16286 AF018169 X61079 S59351 X60142 AF043169
	103241	entrez_X76223 X76223	
	103508	entrez_Y10141 Y10141	•
	103575	entrez_726256 Z26256	
60	119514	NOT_FOUND_entrez_W37937 W37937	
60	121082	genbank_AA398722 AA398722 .	
	128634	AA464918_at	
	105817	genbank_AA397825 AA397825	•
	121518	genbank_AA412155 AA412155	
65	114449	genbank_AA020736 AA020736	•
65	114648	genbank_AA101056 AA101056	
	121950	genbank_AA429515 AA429515	
	107723	genbank_AA015967 AA015967	

Table 3A shows 452 genes up-regulated in chronically diseased lung relative to normal lung. Chronically diseased lung samples represent chronic non-mailgnant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5	Pkey: ExAcon:	E		l identifier number number, Genbank ac	cession number					
	Unigenel Unigene	Title: L	Inigene number Inigene gene title		11 b	00th	af A1 f		-1	
10	R1: R2:	8	oun percentile of Al Oth percentile of Al denocarcinomas	for chronically disease for chronically disease	ed lung samples divided by the ed lung samples divided by the	90th percentile	of normal lu	ma lung sam ng samples, s	pies. quamous cell carc	inomas and
	R3:	7	Oth percentile of Al	for chronically disease ercentile of normal lur	ed lung samples minus the 15t ng samples, squamous celi car	h percentile of A	N for all nom lenocarcinor	nal lung, chron nas minus the	ically diseased lun 15th percentile of	g and tumor samples Al for all normal lung,
15				lung and tumor samp						
	Pkey	ExAccn	UnigenelD	Unigene Title	•	Rí	R2	R3		
20	135423 135378	U50531 AW9618	Hs.138751	Human BRCA2 regi MUM2 protein	ion, mRNA sequence CG030	12.40		2.13		
20	135346	NM_000	928 Hs.992	phospholipase A2,	group IB (pancreas)	12.40		2.10		
	135235 135057	U90268	Hs.93810	ESTs cerebral cavernous		11.67				
25	134951 134799	BE3050 M36821	B1 Hs.169358 Hs.89690	hypothetical protein GRO3 oncogene			8.00 8.20			
	134786	T29618	Hs.89640	TEK tyrosine kinase	e, endothelial (venous	00.00	0.20			
	134772 134752			giutamate receptor, arachidonate 5-lipor	ionotrophic, AMPA 4 xygenase	29.80		1.93		
30	134749 134696	T28499 BE3262	Hs.89485 76 Hs.8861	carbonic anhydrase ESTs	ĬŇ .			2.07		
30	134636	NM_005	582 Hs.87205	lymphocyte antigen	64 (mouse) homolog, r	13.60				
	134627 134622	AI01876 AW9751		glyceronephosphate	e O-acyltransferase ar to A55380 faciogeni			1.92 1.92		
25	134570	U66615	Hs.172280	SWI/SNF related, m	atrix associated, acti	13.20				
35	134561 134468	U76421 NM_001	Hs.85302 772 Hs.83731	CD33 antigen (gp67	se, RNA-specific, B1 (h ')		6.20	1.78		
	134417	NM_006	416 Hs.82921	solute carrier family	35 (CMP-sialic aci					
	134343 134323	D50683 BE17069	Hs.82028 51 : Hs.8700	deleted in liver cand	factor, beta recepto er 1					
40 .	134300 134299	NM_001 AW5809	430 Hs.8136 39 Hs.97199	endothetial PAS dor complement compo						
	134253	X52075	Hs.80738	sialophorin (gpl.115		20.60				
	134182 133985	D52059 L34657	Hs.7972 Hs.78146	KIAA0871 protein platelet/endothelial	cell adhesion molec	12.20				
45	133978	AF0357	l8 Hs.78061	transcription factor 2					•	
•	133835 133651	Al67789 Al30174		RGC32 protein dihydropyrimidinase	-like 2			•		
	133633 133565	D21262 AW9557	Hs.75337 76 Hs.313500		I-body phosphprolein imilar to ALU7_HUMAN A	15.20				
50	133548	ĀW9463	84 Hs.178112	DNA segment, single	e copy probe LNS-CAI/L			1.77		
	133488 133478	AA33529 X83703	95 Hs.74120 Hs.31432	adipose specific 2 cardiac ankyrin repe	eat protein	•	•	2.08		,
	133337	AF08598	33 Hs.293676	ESTs	•		9.60	1.77		
55	133200 133153	AB0377 AF07059		hypothetical protein HSKM-B protein	FLJ 10210	30.60		1.77		
	133130 133120	Al12860 NM_003		zinc finger protein 1	61 ogen-binding protein	22.60				,
	132928	AW1680	82 Hs.169449	protein kinase C, al		13.80				
60	132836 132799	AB02311 W73311		KIAA0960 protein SAC2 (suppressor	of actin mutations 2,	41.60				
	132742	AA0254	30 Hs.292812	ESTs, Weakly simil	ar to T33468 hypotheti	40.40	7.00			
	132548 132476		Hs.193400 I4 Hs.49476	interleukin 6 receptor Homo sapiens clone	or e TUA8 Cri-du-chat regi		7.20 4.76			
65	132439 132240	AK00194 AB0183		hypothetical protein KIAA0781 protein	DKFZp566A1524	21.20		1.88		
05	132210	NM_007	203 Hs.42322	A kinase (PRKA) an	chor protein 2			1.99		
	132199 131751	AL04129 T96555	9 Hs.165084 Hs.31562	ESTs ESTs		15.20		1.76		
70	131745	A182855	9 Hs.31447	ESTs, Moderately s	similar to A46010 X-li	27.80	4.00			
70	131694 131686	NM_000 NM_012		MHC class II transa GRB2-associated b			4.00			
	131676	Al12682 Z45794		ESTs		21.40	6.20			
7.	131629 131589	C18825	Hs.29191	ESTs epithelial membrane	protein 2	21.40				
75	131536 131517			ESTs sema domain, trans	membrane domain (TM),		9.40 3.59			
	131355	R52804	Hs.25956	DKFZP564D206 pro		45.00	4.48			
	131253 131207		Hs.24853 66 Hs.24212	ESTs latrophilin		15.00		1.75		
80	131156	A147220	9 Hs.323117	ESTs			3.54	1.84		
	131066 131061	N64328	Hs.268744	ESTs KIAA1796 protein			J.J4			
	131053 130895				binding protein (G pr DKFZp564L0864 simil	16.60		1.93		
85	130762		Hs.1898	paraoxonase 1		12.00				

	w	O 02/086	443				
	130657	AW337575	Hs.201591	EST ₆			
	130655	AI831962	Hs.17409	cysteine-rich protein 1 (intestinal)			
	130589 130562	AL110226 D50402	Hs.16441 Hs.182611	DKFZP434H204 protein solute carrier family 11 (proton-coupled			2.08 1.91
5	130555	R69743	Hs.116774	integrin, alpha 1		9.60	1.01
	130365	W56119	Hs.155103	eukaryotic translation initiation factor	11.60		
	130273 130259	AW972422 NM_000328	Hs.153863 Hs.153614	MAD (mothers against decapentaplegic, Dr		6.60	1.91
	130090	H97878	Hs.132390	relinitis pigmentosa GTPase regulator zinc finger protein 36 (KOX 18)	21.20		1.51
10	129958	R27498	Hs.1378	annexin A3		5.05	
	129898	AI672731	Hs.13256	ESTs	40.00		
	129875 129699	AA181018 AB007899	Hs.13056 Hs.12017	hypothetical protein FL/13920 homolog of yeast ubiquitin-protein ligas	18.60		
	129626	F13272	Hs.111334	ferritin, light polypeptide			
15	129598	N30436	Hs.11556	Homo sapiens cDNA FLJ12566 fis, clone NT	22.63		
	129593 129565	Al338247 X77777	Hs.98314 Hs.198726	Homo sapiens mRNA; cDNA DKFZp586L0120 (f vasoactive intestinal peptide receptor 1			2.53
	129527	AA769221	Hs.270847	delta-tubulin	39.20		2.00
20	129402	W72062	Hs.11112	ESTs	45.00		2.11
20	129385 129315	AA172106 NM_014563	Hs.110950 Hs.174038	Rag C protein spondyloepiphyseal dysplasia, late	15.20 12.40		
	129312	T97579	Hs.110334	ESTs, Weakly similar to 178885 serine/th	20.83		
	129240	AA361258	Hs.237868	interleukin 7 receptor			1.95
25	129210 129122	AL039940	Hs.202949 Hs.301957	KIAA1102 protein nudix (nucleoside diphosphate linked moi		4.20	
23	129057	AW958473 N90866	Hs.276770	CDW52 antigen (CAMPATH-1 antigen)		4.20	
	128946	Y13153	Hs.107318	kynurenine 3-monooxygenase (kynurenine 3		5.20	
	128798	AF015525	Hs.302043	chemokine (C-C motif) receptor-like 2			2.24
30	128789 128778	AW368576 AA504776	Hs.139851 Hs.186709	caveolin 2 ESTs, Weakly similar to 138022 hypothet	12.20		2.24
50	128766	AW160432	Hs.296460	craniofacial development protein 1	26.40		
		R44238	Hs.155546	KIAA1080 protein; Golgi-associated, gamm			1.78
	128624 128609	BE154765 NM_003616	Hs.102647 Hs.102456	ESTs, Weakly similar to TRHY_HUMAN TRICH survival of motor neuron protein interac	16.00		2.51
35	128603	NM_004915	Hs.102430	ATP-binding cassette, sub-family G (WHIT	12.80		
	128598	AA305407	Hs.102308	potassium inwardly-rectifying channel, s		4.00	
	128458	H55864	Hs.56340	ESTS	47.00		
	128061 127968	AF150882 AA830201	Hs.186877 Hs.124347	sodium channel, voltage-gated, type XII, ESTs	17.20 21.30		
40	127959	Al302471	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L			
	127944	AI557081	Hs.262476	S-adenosylmethionine decarboxylase 1	10.60		
	127925 127896	AA805151 Al669586	Hs.3628 Hs.222194	mitogen-activated protein kinase kinase ESTs	13.40	7.00	
	127859	AA761802	Hs.291559	ESTs	14.00		
45	127817	AA836641	Hs.163085	ESTs	14.00		
	127742 127628	AW293496 Al240102	Hs.180138 Hs.322430	ESTs NDRG family, member 4	11.00 11.10		
	127609	X80031	Hs.530	collagen, type IV, alpha 3 (Goodpasture	11110		
50	127582	AA908954	Hs.130844	ESTs	19.60		
50	127543 127535	AK000787 AA568424	Hs.157392 Hs.164450	Homo saplens cDNA FLJ20780 fis, clone CO ESTs	15.40 17.50		
	127404	AJ379920	Hs.270224	ESTs	14.60		
	127396	L31968	Hs.187991	DKFZP564A122 protein	15.40		,
55	127374 127346	AA442797 AA203616	Hs.312110 Hs.44896	ESTs, Weakly similar to I38022 hypothet DnaJ (Hsp40) homolog, subfamily B, membe	14.60 21.00		
33	127340	BE047653	Hs.119183	ESTs, Weakly similar to ZN91_HUMAN ZINC	15.80		
	127307	AW962712	Hs.126712	ESTs, Wealdy similar to AF191020 1 E2IG5			
	127242 127167	AW390395	Hs.181301	cathepsin S	22.60 21.40		
60	127046	AA625690 AA321948	Hs.190272 Hs.293968	ESTs ESTs	41.20		
-	126928	AA480902	Hs.137401	ESTs	11.00		
	126900	AF137386	Hs.12701	plasmolipin		E 60	1.78
	126852 126816	AA399961 AA248234		gb:zu68c01.r1 Soares_testis_NHT Homo sap gb:csg2228.seq.F Human fetal heart, Lamb	12.20	5.60	
65	126812	AB037860	Hs.173933	nuclear factor I/A	17.19		
	126666	AA648886	Hs.151999	ESTs	13.57		
	126645 126592	AA316181 Al611153	Hs.61635 Hs.6093	six transmembrane epithelial antigen of Homo sapiens cDNA: FLJ22783 fis, clone K	15.40	4.67	
=0	126556	AF255303	Hs.112227	membrane-associated nucleic acid binding	18.00	1.0.	
70	126433	AA325606		gb:EST28707 Cerebellum II Homo sapiens c	16.77		
	126299 126218	AW979155 AL049801	Hs.298275 Hs.13649	amino acid transporter 2 Novel human gene mapping to chomosome 13	14.60	3.50	
	126182	AA721331	Hs.293771	ESTs	13.40	3.30	
75	126177	AW752782	Hs.129750	hypothetical protein FLJ10546	18.20		
75	126142	H86261	Hs.40568	ESTs	14.00		
	126077 125994	M78772 A1990529	Hs.210836 Hs.270799	ESTs ESTs	16.59 17.40		
	125934	AA193325	Hs.32646	hypothetical protein FLJ21901	13.00		
80	125847	AW161885	Hs.249034	ESTs	49.57		
80	125831 125731	H04043 R61771	Hs.26912	gb:yj45c03.r1 Soares placenta Nb2HP Homo ESTs	13.20		
	125676	BE612918	Hs.151973	hypothetical protein FLJ23511	11.20		
	125561	F18572	Hs.22978	ESTs, Weakly similar to ALU4_HUMAN ALU S			,
85	125552	H09701	Hs.278366	ESTs, Weakly similar to 138022 hypotheti	12.60 33.40		
00	125489	H49193	Hs.124984	ESTs, Moderately similar to ALU7_HUMAN A	33.40		

	W	O 02/086	443				
	125422	AA903229	Hs.153717	ESTs			1.80
	125331	AJ422996	Hs.161378	ESTs	38.00		
	125309	T12411	Hs.183745	hypothetical protein FLJ13456	18.20		
_	125167	AL137540	Hs.102541	netrin 4			1.95
5	125139	AW194933	Hs.9788	hypothetical protein MGC10924 similar to	04.00		1.84
	125042	T78906	Hs.269432	ESTs, Moderately similar to ALU1_HUMAN	21.80	10.60	
	124711	NM_004657	Hs.26530	serum deprivation response (phosphatidy)	23.20	10.00	
	124631	NM_014053	Hs.270594	FLVCR protein	21.43		
10	124578 124574	N68321 AL036596	Hs.231500 Hs.42322	EST A kinase (PRKA) anchor protein 2	21.40		1.77
10	124472	N52517	Hs.102670	EST	37.20		••••
	124438	BE178536	Hs.11090	membrane-spanning 4-domains, subfamily A			
	124357	N22401	1101111000	gb:yw37g07.s1 Morton Fetal Cochlea Homo	14.64		
	124306	AW973078	Hs.293039	ESTs		4.00	
15	124214	H58608	Hs.151323	ESTs			
	124097	AW298235	Hs.101689	ESTs		27.20	
	123978	T89832	Hs.170278	ESTs			2.03
	123972	T46848	Hs.70337	Immunoglobulin superfamily, member 4		6.00	4 70
20	123961	AL050184	Hs.21610	DKFZP434B203 protein		45.00	1.79
20	123936	NM_004673	Hs.241519	angiopoletin-like 1		15.80	
	123802	AA620448	11- 040447	gb:ae58c09.s1 Stratagene lung carcinoma		4.23 4.20	
	123734	AA609861	Hs.312447	eSTs gb:no97c02.s1 NCI_CGAP_Pr2 Homo sapiens	33.60	4.20	
	123619 123596	AA602964 AA421130	Hs.112640	EST	10.93		
25	123336	AA384564	Hs.108829	ESTs	10.00		2.18
23	123340	AA504264	Hs.182937	peptidylprolyl isomerase A (cyclophilin	11.20		
	123190	AA489212	Hs.105228	EST	14.20		
	123136	AW451999	Hs.194024	ESTs		7.00	
	123073	AA485061	Hs.105652	ESTs	31.20		,
30	123055	AA482005	Hs.105102	ESTs, Weakly similar to reverse transcri		4.80	
	122699	AA456130	Hs.301721	KIAA1255 protein		5.00	
	122679	AA811286	Hs.192837	ESTs, Weakly similar to ALU5_HUMAN ALU S	14.40		
	122633	NM_001546	Hs.34853	Inhibitor of DNA binding 4, dominant neg	40.00		
25	122553	AA451884	Hs.190121	ESTs	40.00		
35	122544	AW973253	Hs.292689	ESTs	15.40		1.81
	122485	AA524547	Hs.160318 Hs.98849	FXYD domain-containing ion transport reg ESTs, Moderately similar to AF161511 1 H		12.10	1.01
	122211 122127	AA300900 AW207175	Hs.106771	ESTs		12.10	1.95
	122011	AA431082	113.100771	gb:zw78a10.s1 Soares_testis_NHT Homo sap			1.89
40	121992	AI860775	Hs.98506	ESTs		3.60	
	121989	W56487	Hs.193784	Homo sapiens mRNA; cDNA DKFZp586K1922 (f			2.01
	121835	AB033030	Hs.300670	KIAA1204 protein			1.85
	121726	AF241254	Hs.178098	angiotensin I converting enzyme (peptidy	12.43		
	121690	AV660305	Hs.110286	ESTs			1.82
45	121643	AA640987	Hs.193767	ESTs	44.50		
	121633	AA417011	Hs.98175	EST	14.00	40.40	
	121622	AA416931	Hs.126065	ESTs	11.20	16.40	
	121497	AA412031	Hs.97901	EST	12.20		
50	121351	AW206227 W07343	Hs.287727 Hs.182538	hypothetical protein FLJ23132 phospholipid scramblase 4	12.20		1.83
JO	121314 121242	AA400857	Hs.97509	ESTs	22.40		7.00
	121059	AA393283	110.01003	gb:zt74e03.r1 Soares_testis_NHT Homo sap	14.80		
	120934	AA226198		gb:nc26a07.s1 NCI_CGAP_Pr1 Homo sapiens	21.20		
	120755	AA312934	Hs.190745	Homo sapiens cDNA: FLJ21326 fis, clone			1.79
55	120637	AA811804		gb:ob39a05.s1 NCI_CGAP_GCB1 Homo sapiens	20.00		
	120484	AA253170	Hs.96473	EST	40.20		
	120336	N85785	Hs.181165	eukaryotic translation elongation factor		6.60	
	120266	AI807264	Hs.205442	ESTs, Weakly similar to T34036 hypotheti	16.80		
60	120132	W57554	Hs.125019	ESTs		4.73	4 75
60	120041	AA830882	Hs.59368	ESTS		7.20	1,75
	119996 119970	W88996 AA767718	Hs.59134 Hs.93581	EST hypothetical protein FLJ10512	11.20	1.20	
	119861	W78816	Hs.49943	ESTs, Weakly similar to S65657 alpha-1C-	11.20	3.78	
	119824		Hs.184	advanced glycosylation end product-speci		2	
65	119740	AW021407	Hs.21068	hypothetical protein	20.20		
	119271	AI061118	Hs.65328	Fanconi anemia, complementation group F	15.20		
	119221	C14322	Hs.250700	tryptase beta 1			
	119126	R45175	Hs.117183	ÉSTs	12.60		
	119073	BE245360	Hs.279477	ESTs			
70	118928	AA312799	Hs.283689	activator of CREM in testis		10.00	
	118901	AW292577	Hs.94445	ESTs		3.96	
	118661	AL137554	Hs.49927	protein kinase NYD-SP15	40.40	9.60	
	118607	AI377444 ·	Hs.54245	ESTs, Weakly similar to S65824 reverse t	10.40		1.90
75	118449	A1813865	Hs.164478	hypothetical protein FLJ21939 similar to FKBP-associated protein	16.20		1.30
, 5	118416 118379	N66028 N64491	Hs.49105 Hs.48990	ESTs	10.20	4.00	
	118329	N63520	Hs.48990	gb:yy62f01.s1 Soares_multiple_sclerosis_		6.60	
	118329	N63451	Hs.141600	ESTs, Weakly similar to alternatively s		3.80	
	118253	AA497044	Hs.20887	hypothetical protein FLJ10392	17.60		
80	118124	N56968	Hs.46707	chromosome 21 open reading frame 37	14.00		
	118056	AB037746	Hs.42768	hypothetical protein DKFZp76100113			1.86
	118032	N52802	Hs.47544	EST		5.00	
	117840	T26379	Hs.48802	Homo sapiens clone 23632 mRNA sequence		4.00	4 00
95	117404	N39725	Hs.15220	zinc finger protein 106	14.00		1.90
85	117314	N32498	Hs.42829	ESTs	14.20		

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	W	O 02/086	443				
	117209	W03011	Hs.306881	MSTP043 protein			
	117023	AW070211	Hs.102415	Homo sapiens mRNA; cDNA DKFZp586N0121 (f	20.20		2.31
	116814 116784	H50834 AB007979	Hs.301281	gb:yp86a10.s1 Soares fetal liver spleen Homo sapiens mRNA, chromosome 1 specific	20.20	3.51	
5	116766	Al608657	Hs.95097	ESTs	16.20		
	116712	AW901618	Hs.61935	Homo sepiens mRNA; cDNA DKFZp7611071 (fr	10.00	6.80	
	116707 116351	H10344 AL133623	Hs.49050 Hs.82501	ESTs, Wealty similar to A Chain A, Human similar to mouse Xm1 / Dhm2 protein	18.60 19.40		
10	116279	AW971248	Hs.291289	ESTs, Weakly similar to ALU1_HUMAN ALU S			
10	116166	AL039940	Hs.202949	KIAA1102 protein			2.13
	116152 116117	AL040521 BE613410	Hs.15220 Hs.31575	zinc finger protein 106 SEC63, endoplasmic reticulum translocon	13.20		1.75
	116107	AL133916	Hs.172572	hypothetical protein FLJ20093	30.11		
1.5	115965	AA001732	Hs.173233	hypothetical protein FLJ10970			2.36
15	115955 115844	AF263613 Al373062	Hs.44198 Hs.332938	Intracellular membrane-associated calciu hypothetical protein MGC5370	18.20 18.57		
	115683	AF255910	Hs.54650	Junctional adhesion molecule 2	10.01	23.00	
•	115673	AA406341	Hs.269908	Homo saplens cDNA FLJ11991 fis, clone HE	11.82		
20	115672	Al889110	Hs.73251	ESTs	10.60		4.70
20	115566 115313	AI142336 AA808001	Hs.43977 Hs.184411	Human DNA sequence from clone RP11-196N1 albumin	25.20		1.76
	115279	AW964897	Hs.290825	ESTs		8.00	
	115230	AA278300	Hs.124292	Homo saplens cDNA: FLJ23123 fis, clone L	44.00		1.80
25	115110 114999	AK001671 BE246481	Hs.11387 Hs.87856	KIAA1453 protein ESTs	14.20 19.20		
23	114930	AA237022	Hs.188717	ESTs	13.20	5.60	
	114922	AA235672	Hs.87491	ESTs		3.60	
	114837	BE244930	Hs.166895	ESTs	43.70 11.00		
30	114769 114761	AA149060 AA143781	Hs.296100 Hs.126280	ESTs hypothetical protein FLJ23393	14.00		
-	114736	Al610347	Hs.103812	ESTs, Moderately similar to ALU1_HUMAN A		4.20	
	114596	AA310162	Hs.169248	cytochrome c	10.71		
	114518 114455	AW163267 H37908	Hs.106469 Hs.271616	suppressor of var1 (S.cerevisiae) 3-like ESTs, Weakly similar to ALU8_HUMAN ALU S	20.40 20.40		
35	114452	Al369275	Hs.243010	Homo saplens cDNA FLJ14445 fis, clone HE	20.40	17.20	
	114359	NM_016929	Hs.283021	chloride intracellular channel 5			2.09
	114357	R41677	Hs.6107	Homo sapiens cDNA FLJ14839 fis, clone OV	12.40		2.00
	114251 114138	H15261 AW384793	Hs.21948 Hs.15740	ESTs Homo sapiens mRNA; cDNA DKFZp434E033 (fr		11.40	2.00
40	114124	W57554	Hs.125019	ESTs		6.04	
	113946	AW083883	Hs.37896	Homo sapiens cDNA FLJ13510 fis, clone PL			1.82
	113695 113606	T96965 NM_013343	Hs.17948 Hs.278951	ESTs, Weakly similar to ALUB_HUMAN !!!! NAG-7 protein			2.15
	113590	R49642	Hs.142447	ESTs, Weakly similar to ALU1_HUMAN ALU S		3.60	2
45	113560	T91015	Hs.268626	ESTs	32.00		
	113552 113540	AI654223 AW152618	Hs.16026 Hs.16757	hypothetical protein FLJ23191 ESTs			
	113502	T89130	115.107.57	gb:ye12d01.s1 Stratagene lung (937210) H		8.35	
50	113288	A1076838	Hs.12967	ESTs	12.40		
50	113252	NM_004469	Hs.11392	c-fos induced growth factor (vascular en		4.27	
	113238 113203	R45467 AA743563	Hs.189813 Hs.10305	ESTs ESTs	21.20		. 4
	113195	H83265	Hs.8881	ESTs, Weakly similar to S41044 chromosom			1.92
55	113089	T40707	Hs.270862	ESTs	14.33	. 0.00	
33	113076 113009	AF033199 T23699	Hs.8198 Hs.7246	zinc finger protein 204 ESTs		· 6.00 9.40	
	112937	AI694320	Hs.6295	ESTs, Weakly similar to T17248 hypotheti		12.20	
	112891	T03927	Hs.293147	ESTs, Moderately similar to A46010 X-li	10.57		
60	112794 112691	R97018 R88708	Hs.220647	gb:yq74b08.s1 Soares fetal liver spleen ESTs	26.60 15.33		
00	112602	AW004045	Hs.203365	ESTs	15.60		
	112366	AF035318	Hs.12533	Homo sapiens clone 23705 mRNA sequence	15.40		
	112210 112064	R49645 AL049390	Hs.7004 Hs.22689	ESTs Homo saplens mRNA; cDNA DKFZp586O1318 (f	14.00 13.00		
65	111998	R42379	Hs.138283	ESTs	11.00		
	111987	NM_015310	Hs.6763	KIAA0942 protein	22.40		
	111803	AA593731	Hs.325823	ESTs, Moderately similar to ALU5_HUMAN A			1.77
	111737 111605	H04607 T91061	Hs.9218 Hs.194178	ESTs ESTs, Moderately similar to PC4259 ferri	23.00		1.86
70	111510	R07856	Hs.16355	ESTs	11.02		
	111341	AL157484	Hs.22483	Homo sapiens mRNA; cDNA DKFZp762M127 (fr	40.40		1.88
	111280 111247	AA373527 AW058350	Hs.19385 Hs.16762	CGI-58 protein Homo sapiens mRNA; cDNA DKFZp564B2062 (f	18.40		
	111232	Al247763	Hs.16928	ESTs	27.60		
75	110942	R63503	Hs.28419	ESTs	14.80		
	110924 110837	AW058463 H03109	Hs.12940 Hs.108920	zinc-fingers and homeoboxes 1 HT018 protein	24.71		2.18
	110824	Al767183	Hs.26942	ESTs	12.20		2.10
90	110776	AB032417	Hs.19545	frizzled (Drosophila) homolog 4			1.75
80	110576 110369	H60869 AK000768	Hs.37889	ESTs	13.00	5.60	
	110099	R44557	Hs.107872 Hs.23748	hypothetical protein FLJ20761 ESTs		J.00	2.31
	109984	AI796320	Hs.10299	Homo sapiens cDNA FLJ13545 fis, clone PL			
85	109958	AA001266	Hs.133521	ESTs	11.25		2 60
$\sigma_{\mathcal{J}}$	109893	AAB84208	Hs.30484	ESTs			2.68

Homo sapiens mRNA; cDNA DKFZp434M229 (fr

succinate-CoA ligase, GDP-forming, alpha

CD83 antigen (activated B lymphocytes, i

RAS p21 protein activator (GTPase activa

hypothetical protein FLJ10849

caveolin 1, caveolae protein, 22kD

flavin containing monooxygenase 2

A kinase (PRKA) anchor protein 1

104074

103749

103645

103554

103541

103496

103428

103353

80

85

AL162039

AL135301

AW246253

A1878826

AI815601

BE383507

Y09267

X89399

Hs.31422

Hs.8768

Hs.7043

Hs.323469

Hs.79197

Hs.132821

Hs.78921

Hs.119274

11.20

10.86

12.00

11.20

19.80

· 1.80

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	103295	X81479	Hs.2375	egf-like module containing, mucin-like,		3.60	
	103280	U84722	Hs.76206	cadherin 5, type 2, VE-cadherin (vascula			1.76
	103100	NM_005574	Hs.184585	LIM domain only 2 (rhombotin-like 1)			2.15
5	103025	NM_002837	Hs.123641	protein tyrosine phosphatase, receptor t			2.13
)	102698	M18667	Hs.1867	progastricsin (pepsinogen C)	11.00		
	102659 102580	BE245169 U60808	Hs.211610 Hs.152981	CUG triplet repeat, RNA-binding protein CDP-diacylglycerol synthase (phosphatida	25.40		
	102560	AA034127	Hs.153487	signal transducing adaptor molecule (SH3	14.00		
	102363	NM_003734	Hs.198241	amine oxidase, copper containing 3 (vasc			
10	102302	AA306342	Hs.69171	protein kinase C-like 2	10.86		
10	102283		Hs.83381	quanine nucleotide binding protein 11			
	102188	U20350	Hs.78913	chemokine (C-X3-C) receptor 1		7.40	
	102151	T27013	Hs.3132	steroidogenic acute regulatory protein	16.40		
	101957	L28824	Hs.74101	spleen tyrosine kinase	15.40		
15	101842	M93221	Hs.75182	mannose receptor, C type 1			
	101771	NM_002432	Hs.153837	myeloid cell nuclear differentiation ant			4 70
	101764	Al198550	Hs.81256	S100 calcium-binding protein A4 (calcium	18.80		1.78
	101716	AF050658	Hs.2563	tachykinin, precursor 1 (substance K, su	10.00		2.22
20	101678	M62505	Hs.2161	complement component 5 receptor 1 (C5a l	504.80		2.46
20	101447	M21305	Hs.79345	gb:Human alpha satellite and satellite 3 coagulation factor VIII, procoagulant co	304.00	31.00	
	101383 101346	NM_000132 Al738616	Hs.77348	hydroxyprostaglandin dehydrogenase 15-(N		01.00	1.75
	101346	NM 005795	Hs.152175	calcitonin receptor-like		•	•
	101336	NM_006732	Hs.75678	FBJ murine osteosarcoma viral oncogene h			2.24
25	101330		Hs.80261	enhancer of filamentation 1 (cas-like do			
	101277	BE297626	Hs.296049	microfibrillar-associated protein 4			
	101262			gb:Human dystrophin (dp140) mRNA, 5' end	19.00		
	101168	NM_005308	Hs.211569	G protein-coupled receptor kinase 5			2.01
	101102	NM_003243	Hs.79059	transforming growth factor, beta recepto			
30	101088	X70697	Hs.553	solute carrier family 6 (neurotransmitte		7.52	
	101066	AW970254	Hs.889	Charot-Leyden crystal protein	19.38		1.91
	100971	BE379727	Hs.83213	fatty acid binding protein 4, adipocyte	15.40		1.91
	100893	BE245294	Hs.180789	S164 protein	11.20		
35	100770			amyloid beta (A4) precursor protein (pro	14.80		
22	100716 100555	X89887 M69181	Hs.172350	HIR (histone cell cycle regulation defec gb:Human nonmuscle myosin heavy chain-B	33.00		
	100555	NM_014747	Hs.78748	KIAA0237 gene product	16.20		•
	100423	D86640	Hs.56045	src homology three (SH3) and cysteine ri	10.20	4.00	•
	100382	D83407	Hs.156007	Down syndrome critical region gene 1-lik		4.24	
40	100351	D64158				6.20	
. •	100299	D49493	Hs.2171	growth differentiation factor 10		21.20	
	100134	AA305746	Hs.49	macrophage scavenger receptor 1			
	100108	U09577	Hs.76873	hyaluronoglucosaminidase 2			1.79
	100095	Z97171	Hs.78454	myodilin, trabecular meshwork inducible	44.00	5.40	
45	100066			•	11.29		
				the state of the s			

TABLE 3B shows the accession numbers for those primekeys tacking unigenelD's for Table 3A. For each probeset we have listed the gene cluster number from which the oilgonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

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Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers

85

60	Pkey	CAT number	Accession	s	
00	123619	371681_1	AA602964	AA609200	
	126433	127143_1	AA325606	AA099517 N89423	
	125831	1522905 1	H04043 D	60988 D60337	
	126816			AA090985	
65	126852			AA128347	
••	121059			AA398628	
	120637	200885_1		AA809404 AA286907 AW	977624
	122011		AA43108		
	120934	177521 1	AA226198	AA226513 AA383773	
70	123802			AA620448	
	116814	genbank_H508		H50834	
	118329		20	N63520	
	104404				
	104776			AA026349	
75	113502				
	101262				
	108573			AA086005	
•	101447				
	124357			N22401	
80	108781			AA128654	
	112794			R97018	
	100351				
	100555	tiar HT2245		A81105 U51039	

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Table 4A shows 202 genes up-regulated in samples from patients treated with chemotherapy or radiotherapy. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

Pkey: Unique Eos probeset identifier number
ExAccn: Exemplar Accession number, Genbank accession number
UnigenelD: Unigene number
Unigene Title: Unigene gene title 5

	Unigene	Title: Unigene g	ene title		
	R1:	average of	Al for samples	s from patients treated with chemotherapy or radiother	apy divided by the average of Al for normal lung samples.
10		•			
	Pkey	ExAcen	UnigenelD	Unigene Title	R1
	100113	NM_001269	Hs.84746	chromosome condensation 1	27.20
1 ~	100187	D17793	Hs.78183	aldo-keto reductase family 1, member C3	20.60
15	100210	D26361	Hs.3104	KIAA0042 gene product	. 20.40
	100225	D28539	Hs.167185	glutamate receptor, metabotropic 5	20.60
	100269	NM_001949	Hs.1189	E2F transcription factor 3	29.40
	100438	AA013051	Hs.91417	topoisomerase (DNA) II binding protein	23.50
20	100877	X80821	Hs.27973	KIAA0874 protein	35.56 43.40
20	100893	BE245294	Hs.180789	S164 protein	21.80
	101273	Z11933	Hs.182505	POU domain, class 3, transcription facto	193.60
	101447	M21305	11- 4000	gb:Human alpha satellite and satellite 3	38.40
	101649	AW959908	Hs.1690 Hs.620	heparin-binding growth factor binding pr bullous pemphigoid antigen 1 (230/240kD)	198.80
25	101724 101748	L11690 NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	78.60
23	101748	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	162.20
	101879	AA176374	Hs.243886	nuclear autoantigenic sperm protein (his	50.00
	101915	AF207881	Hs.155185	cytosolic ovarian carcinoma antigen 1	26.00
	101973	U41514	Hs.80120	UDP-N-acetyl-alpha-D-galactosamine:polyp	37.20
30	102025	U04045	Hs.78934	mutS (E. coli) homolog 2 (colon cancer,	•
-	102031	U04898	Hs.2156	RAR-related orphan receptor A	32.00
	102052	NM_002202	Hs.505	ISL1 transcription factor, LIM/homeodoma	51.20
	102391	AA296874	Hs.77494	deoxyguanosine kinase	13.90
	102420	U44060	Hs.14427	Homo sapiens cDNA: FLJ21800 fis, clone H	28.80
35	102610	U65011	Hs.30743	preferentially expressed antigen in mela	110.60
	102829	NM_006183	Hs.80962	neurotensin	116.80
	103000	NM_001975	Hs.146580	enolase 2, (gamma, neuronal)	2.30
	103036	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial	181.40
	103507	AJ000512	Hs.296323	serum/glucocorticoid regulated kinase	49.20
40	103587	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein	86.60
	104660	BE298665	Hs.14846	Homo sapiens mRNA; cDNA DKFZp564D016 (fr	42.60
•	104896	AW015318	Hs.23165	ESTs	29.40
	105038	AW503733	Hs.9414	KIAA1488 protein	21.50
45	105298	BE387790	Hs.26369	hypothetical protein FLJ20287	32.80
45	105510	Z42047	Hs.283978	Homo sapiens PRO2751 mRNA, complete cds	20.20
	105667	AA767526	Hs.22030	paired box gene 5 (B-cell lineage specif	28.40
	106073	AL157441	Hs.17834	downstream neighbor of SON	25.40 32.00
	106205	AW965058	Hs.111583	ESTs, Weakly similar to I38022 hypotheti	40.60
50	106516	AL137311	Hs.234074	Homo sapiens mRNA; cDNA DKFZp761G02121 (59.80
30	106533	AL134708	Hs.145998	ESTs ESTs	43.40
	106575 106654	AW970602 AW075485	Hs.105421 Hs.286049	phosphoserine aminotransferase	50.80
	106851	A1458623	113.200043	gb:tk04g09.x1 NCI_CGAP_Lu24 Homo sapiens	53.40
	106995	AB023139	Hs.37892	KIAA0922 protein	20.88
55	107332	T87750	Hs.183297	DKFZP566F2124 protein	23.60
-	107532		Hs.173684	Homo sapiens mRNA; cDNA DKFZp762G207 (fr	57.20
	107922	BE153855	Hs.61460	lg superfamily receptor LNIR	49.00
	108609	BE409857	Hs.69499	hypothetical protein	19.67
	108780	AU076442	Hs.117938	collagen, type XVII, alpha 1	48.17
60	109166	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines	59.20
	109260	AW978515	Hs.131915	KIAA0863 protein	28.60
٠.	109280	AK001355	Hs.279610	hypothetical protein FLJ10493	22.80
	109292	AW975746	Hs.188662	KIAA1702 protein	1
~~	109384	AA219172	Hs.86849	ESTs	21.00
65	109415	U80736	Hs.110826	trinucleotide repeat containing 9	31.60
•	109445	AA232103	Hs.189915	ESTs	24.20
		AW967069	Hs.211556	hypothetical protein MGC5487	21.40
	109633	AW003785	Hs.170267	ESTs	20.40
70	109786	Al989482	Hs.146286	kinesin family member 13A	19.60
70 ·	109958	AA001266	Hs.133521	ESTs	24.00 28.40
	110920	N47224	Hs.20521	HMT1 (hnRNP methyltransferase, S. cerevi	36.00
	110924	AW058463	Hs.12940	zinc-fingers and homeoboxes 1	61.20
	111084	H44186	Hs.15456	PDZ domain containing 1	24.60
75	111132		Hs.83293	hypothetical protein	27.20
15	111229	AW389845 AA837396	Hs.110855	ESTs LIS1-interacting protein NUDE1, rat homo	48.00
	111337 111987	NM_015310	Hs.263925 Hs.6763	KIAA0942 protein	37.80
	112046	AA383343	Hs.22116	CDC14 (cell division cycle 14, S. cerevi	26.80
	112268	W39609	Hs.22003	solute carrier family 6 (neurotransmitte	63.80
80	112685	R87650	Hs.33439	ESTs, Weakly similar to ALU1_HUMAN ALU	26.40
	112871	AL110216	Hs.12285	ESTs, Weakly similar to 155214 salivary	47.64
	112897	AW206453	Hs.3782	ESTs	22.00
	112973	AB033023	Hs.318127	hypothetical protein FLJ 10201	65.00
	112992	AL157425	Hs.133315	Homo sapiens mRNA; cDNA DKFZp761J1324 (f	42.00
85	113073	N39342	Hs.103042	microtubule-associated protein 18	55.40
			· -	•	

	M	/ O 02/086	443		
	113494	T91451	Hs.86538	ESTs	22.80
	113560	T91015	Hs.268626	ESTs	22.80
	113849		Hs.8858	bromodomain adjacent to zinc finger doma	51.80
_	113950		Hs.30504	Homo sapiens mRNA; cDNA DKFZp434E082 (fr	28.20
5	114339		Hs.22790	ESTs	20.20
	114365		Hs.18653	hypothetical protein FLJ14627	21.00
	114455	H37908	Hs.271616	ESTs, Weakly similar to ALU8_HUMAN ALU S	25.80
	114518	AW163267	Hs.106469	suppressor of var1 (S.cerevisiae) 3-like	23.60
10	114824	AA960961	Hs.305953	zinc finger protein 83 (HPF1)	27.20
10	114837	BE244930	Hs.166895	ESTs	30.20
	114974	AW966931	Hs.179662	nucleosome assembly protein 1-like 1	20.80
	115075	AA814043	Hs.88045	ESTs	30.60
	115084	BE383668	Hs.42484	hypothetical protein FLJ10618	28.86
1.5	115291	BE545072	Hs.122579	hypothetical protein FLJ10461	38.00
15	115313	AA808001	Hs.184411	albumin	22.60
	115697	D31382	Hs.63325	transmembrane protease, serine 4	173.60
	115909	AW872527	Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH	27.77
	116090	AI591147	Hs.61232	ESTs	20.80
20	116107	AL133916	Hs.172572	hypothetical protein FLJ20093	164.20
20	116399	AA889120	Hs.110637	homeo box A10	38.00
	117099	H93699		gb:yv16a11.s1 Soares fetal liver spleen	21.60
	117881	AF161470	Hs.260622	butyrate-induced transcript 1	49.40
	118091	AW005054	Hs.47883	ESTs, Wealdy similar to KCC1_HUMAN CALCI	22.40
25	118138	AA374756	Hs.93560	Homo sapiens mRNA for KIAA1771 protein,	22.00
23	118720	N73515	11- 44633	gb:za49d07.s1 Soares fetal liver spleen	20.00
	118873	AI824009	Hs.44577	ESTs	19.40
	119126	R45175	Hs.117183	ESTs	111.20
	119717	AA918317	Hs.57987	B-cell CLL/lymphoma 11B (zinc finger pro	33.00
30	119940	AL050097	Hs.272531	DKFZP586B0319 protein	31.00
20	120266	AI807264	Hs.205442	ESTs, Weakly similar to T34036 hypotheti	20.20
	120515	AA258356	Un 4040	gb:zr59c10.s1 Soares_NhHMPu_S1 Homo sapi	25.00
	120859 120983	AA826434 AA398209	Hs.1619	achaete-scute complex (Drosophila) homol	95.40 105.20
	121054	AW976570	Hs.97587	EST ESTs	
35	121369	AW450737	Hs.97387 Hs.128791		38.80 41.60
55	122335	AA443258	Hs.241551	CGI-09 protein chloride channel, calcium activated, fam	30.80
	122612	AA974832	Hs.128708	ESTs	19.60
	123130	AA487200	113.120700	gb:ab19f02.s1 Stratagene lung (937210) H	33.20
	123440	A1733692	Hs.112488	ESTs	23.17
40	123596	AA421130	Hs.112640	EST	23.00
10	123619	AA602964	110.112070	gb:no97c02.s1 NCI_CGAP_Pr2 Homo sapiens	28.80
	124006	Al147155	Hs.270016	ESTs .	77.60
	124169	BE079334	Hs.271630	ESTs	22.20
	124281	Al333756	Hs.111801	arsenate resistance protein ARS2	42.20
45	124472	N52517	Hs.102670	EST	32.60
	124617	AW628168	Hs.152684	ESTs	21.80
	124631	NM_014053	Hs.270594	FLVCR protein	30.40
	124839	R55784	Hs.140942	ESTs	21.20
	125186	AA610620	Hs.181244	major histocompatibility complex, class	42.80
50	125321	T86652	Hs.178294	ESTs	27.00
	125535	NM_013243	Hs.22215	secretogranin III	23.80
	125646	AA628962	Hs.75209	protein kinase (cAMP-dependent, catalyti	23.20
	125684	AW589427	Hs.158849	Homo sapiens cDNA: FLJ21663 fis, clone C	21.20
	125724	AL360190	Hs.295978	Homo sapiens mRNA full length insert cDN	48.80
55	125847	AW161885	Hs.249034 *	ESTs	31.00
	125934	AA193325	Hs.32646	hypothetical protein FLJ21901	21.20
	126077	M78772	Hs.210836	ESTs	49.80
-	126299	AW979155	Hs.298275	arnino acid transporter 2	21.80
	126395	A1468004	Hs.278956	hypothetical protein FLJ12929	71.00
60	126433	AA325606		gb:EST28707 Cerebellum II Homo sapiens c	23.20
	126509	R47400	Hs.23850	ESTs	23.80
	126538	AB030656	Hs.17377	coronin, actin-binding protein, 1C	23.10
	126666	AA648886	Hs.151999	ESTs	36.00
~~	126812	AB037860	Hs.173933	nuclear factor I/A	20.80
65	126872	AW450979		gb:UI-H-BI3-ala-a-12-0-UI.s1 NCI_CGAP_Su	46.29
	127046	AA321948	Hs.293968	ESTs	22.80
	127431	AW771958	Hs.175437	ESTs, Moderately similar to PC4259 ferri	30.00
	127489	AA650250	Hs.272076	ESTs	20.80
70	127521	AW297206	Hs.164018	ESTs	25.20
70	127742	AW293496	Hs.180138	ESTs	28.00
	127925	AA805151	Hs.3628	mitogen-activated protein kinase kinase	21.20
	127930	AA809672	Hs.123304	ESTs	20.54
	127968	AA830201	Hs.124347	ESTs	28.20
75	127987	Al022103	Hs.124511	ESTs	19.60
75	128116	H07103	Hs.286014	Homo sapiens, clone IMAGE:3867243, mRNA	20.40
	128609	NM_003616	Hs.102456	survival of motor neuron protein interac	34.40
	128777	Al878918	Hs.10526	cysteine and glycine-rich protein 2	53.80
	128949	AA009647	Hs.8850	a disintegrin and metalloproteinase doma	23.00
90	129168	A1132988	Hs.109052	chromosome 14 open reading frame 2	37.60
80	129404	AJ267700	Hs.317584	ESTs	28.60
	129527	AA769221	Hs.270847	delta-tubulin .	40.80
	129574	AA026815	Hs.11463	UMP-CMP kinase	31.20
	129598	N30436	Hs.11556	Homo sapiens cDNA FLJ12566 fis, clone NT	29.60
85	129785	H19006	Hs.184780	ESTs	72.20
J	129970	AV655806	Hs.296198	chromosome 12 open reading frame 4	22.20

	W	O 02/086	443		
	130149	AW067805	Hs.172665	methylenetetrahydrofolate dehydrogenase	29.60
	130199	Z48579	Hs.172028	a disintegrin and metalloproteinase doma	27.60
	130441	U63630	Hs.155637	protein kinase, DNA-activated, catalytic	28.36
	130466	W19744	Hs.180059	Homo sapiens cDNA FLJ20653 fis, clone KA	20.20
5	130482	AW409701	Hs.1578	bacutoviral IAP repeat-containing 5 (sur	22.40
_	130617	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	19.60
	130703	R77776	Hs.18103	ESTs	19.40
	130732	AW890487	Hs.63984	cadherin 13, H-cadherin (heart)	21.40
	130867	NM_001072	Hs.284239	UDP glycosyltransferase 1 family, polype	110.00
10	131028	Al879165	Hs.2227	CCAAT/enhancer binding protein (C/EBP),	25.20
	131086	AL035461	Hs.2281	chromogranin B (secretogranin 1)	40.60
	131284	NM_001429	Hs.25272	E1A binding protein p300	24.60
	131775	AB014548	Hs.31921	KIAA0648 protein	21.00
	131860	BE383676	Hs.334	Rho guanine nucleotide exchange factor (33.40
15	131945	NM_002916	Hs.35120	replication factor C (activator 1) 4 (37	60.80
	132040	NM_001196	Hs.315689	Homo sapiens cDNA: FLJ22373 fis, clone H	. 20.40
	132084	NM_002267	Hs.3886	karyopherin alpha 3 (importin alpha 4)	29.40
	132389	AA310393	Hs.190044	ESTs	32.40
	132437	AA152106	Hs.4859	cyclin L ania-6a	27.40
20	132550	AW969253	Hs.170195	bone morphogenetic protein 7 (osteogenic	75.60
	132617	AF037335	Hs.5338	carbonic anhydrase XII	31.36
	132632	AU076916	Hs.5398	guanine monphosphate synthetase	32.40
	132672	W27721	Hs.54697	Cdc42 guanine exchange factor (GEF) 9	23.40
	132742	AA025480	Hs.292812	ESTs, Weakly similar to T33468 hypotheti	61.20
25	132771	Y10275	Hs.56407	phosphoserine phosphatase	22.33
	133070	U92649	Hs.64311	a disintegrin and metalloproteinase doma	23.50
	133153	AF070592	Hs.66170	HSKM-B protein	30.00
	133181	X91662	Hs.66744	twist (Drosophila) homolog (acrocephalos	23.80
	133282	AA449015	Hs.286145	SRB7 (suppressor of RNA polymerase B, ye	51.60
30	133350	Al499220	Hs.71573	hypothetical protein FLJ10074	33.00
	133592	AV652066	Hs.75113	general transcription factor IIIA	82.00
	133658	AA319146	Hs.75426	secretogranin II (chromogranin C)	
	133865	AB011155	Hs.170290	discs, large (Drosophila) homolog 5	69.33
	134032	NM_005025	Hs.78589	serine (or cysteine) proteinase inhibito	33.20
35	13/125	NM 014781	He 50421	KIAA0203 nene product	31.60

serine (or cysteine) proteinase inhibito KIAA0203 gene product BCL2/adenovirus E1B 19kD-interacting pro

ESTs, Moderately similar to A46010 X-lin

phosphoribosylglycinamide formyltransfer SWI/SNF related, matrix associated, acti

dual-specificity tyrosine-(Y)-phosphoryl

hydroxysteroid (17-beta) dehydrogenase

cyclin-dependent kinase 5, regulatory su

Gantigen 7B

neurotrophin 3

AA487200

TABLE 4B shows the accession numbers for those primekeys tacking unigenelD's for Table 4A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

31.60

30.60

23.40

49.20

20.20

20.80

37.60

53.40

31.60

28.80

PCT/US02/12476

Pkey: Unique Eos probeset CAT number: Gene cluster number Unique Eos probeset identifier number Accession: Genbank accession numbers

CAT number Accessions Pkey 123619 371681_1 AA602964 AA609200 60 AA325606 AA099517 N89423 126433 127143_1 AW450979 AA136653 AA136656 AW419381 AA984358 AA492073 BE168945 AA809054 AW238038 BE011212 BE011359 BE011367 BE011368 BE011368 BE011215 BE011365 BE011363 126872 142696_1 106851 322947_1 AI458623 AA639708 AA485409 R22065 AA485570 118720 genbank_N73515 N73515 65 120515 genbank_AA258356 AA258356 117099 321871_1 H93699 H97976 H80036

entrez_M21305 M21305

genbank_AA487200

NM_005025 NM_014781

U15174

BE538082

AA339449

NM_006482 AA448542

U66615

H58818

X53655

AL134197

134753

135002

135029

135047

135345

Hs.50421

Hs.79428

Hs.8172

Hs.82285

Hs.172280

Hs.173135

Hs.251677

Hs.187579

Hs.93597

Hs.99171

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101447 123130

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PCT/US02/12476
Table 5A shows 680 genes up-regulated in squamous cell carcinoma or adenocarcinoma lung tumors relative to normal lung and chronically diseased lung. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

	5	Pkey:	Unique Eos probeset identifier number
•		ExAcon:	Exemplar Accession number, Genbank accession number
		UnigenelD:	Unigene number
		Unigene Title:	Unigene gene tille
	_	R1:	70th percentile of AI for squamous cell carcinoma and adenocarcinoma lung tumor samples divided by the 90th percentile of AI for normal and chronically
- 1	0		diseased lung samples.
		R2:	80th percentile of Al adenocarcinoma lung tumor samples divided by the 90th percentile of Al for normal and chronically diseased lung samples.
		R3:	80th percentile of Al squamous cell carcinoma lung tumor samples divided by the 90th percentile of Al for normal and chronically diseased lung samples.
		R4:	80th percentile of Al adenocarcinoma lung tumor samples divided by the 80th percentile of Al for squamous cell carcinoma lung tumor samples.
_	_	R5:	70th percentile of Al for squamous cell carcinoma and adenocarcinoma lung tumor samples minus the 15th percentile of Al for all normal lung, chronically
1	5		diseased lung and tumor samples divided by 90th percentile of AI for normal and chronically diseased lung samples minus the 15th percentile of AI for all
_	_		normal lung, chronically diseased lung and tumor samples
1	5		70th percentile of Al for squamous cell carcinoma and adenocarcinoma lung tumor samples minus the 15th percentile of Al for all normal lung, chronically diseased lung and tumor samples divided by 90th percentile of Al for normal and chronically diseased lung samples minus the 15th percentile of Al for all

		110.111		,					•
	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2	R3	R4	R5
20	,								
	100035			AFFX control: GAPDH					6.76
	100036			AFFX control: GAPDH					5.77 5.75
	100037	100100		AFFX control: GAPDH		8.00			5.75
25	100071	A28102	He pages	Human GABAa receptor alpha-3 subunit		0.00			5.71
23	100114 100154	X02308 H60720	Hs.82962 Hs.81892	thymidylate synthetase KIAA0101 gene product	3.84				0
	100187	D17793	Hs.78183	aldo-keto reductase family 1, member C3	3.33				
	100188	AW247090	Hs.57101	minichromosome maintenance deficient (S.					4.52
	100202	BE294407	Hs.99910	phosphofructokinase, platelet					5.49
30	100216	AA489908	Hs.1390	proteasome (prosome, macropain) subunit,					5.67
	100269	NM_001949	Hs.1189	E2F transcription factor 3	2.55				
	100287	AU076657	Hs.1600	chaperonin containing TCP1, subunit 5 (e					5.66
	100297	AU077258	Hs.182429	protein disulfide isomerase-related prot					3.81
25	100330	AW410976	Hs.77152	minichromosome maintenance deficient (S.	r 07				4.50
35	100335	AW247529	Hs.6793	platelet-activating factor acetylhydrola	5.07				4.82
	100360	W70171	Hs.75939	uridine monophosphate kinase					3.79
	100372	NM_014791	Hs.184339	KIAA0175 gene product			•	15.65	0.73
	100474 100486	NM_000699 T19006	Hs.300280 Hs.10842	amylase, alpha 2A; pancreatic RAN, member RAS oncogene family				10.00	5.49
40	100400	D56165	Hs.275163	non-metastatic cells 2, protein (NM23B)					4,17
40	100516	D90278	Hs.11	carcinoembryonic antigen-related cell ad		7.20			
	100522	X51501	Hs.99949	prolactin-induced protein				14.20	
	100559	NM_000094	Hs.1640	collagen, type VII, alpha 1 (epidermolys	3.10				
	100576	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid				9.30	
45	100629	AA015693	Hs.21291	mitogen-activated protein kinase kinase				20.60	
	100661	BE623001	Hs.132748	Homo sapiens ribosomal protein L39 mRNA,	3.85				
		- AA353686	Hs.57813	zinc ribbon domain containing, 1		8.60		10.00	
	100696	D14887	Hs.121686	general transcription factor IIA, 1 (37k			24.80	10.00	
50	100709	N26539	Hs.100469	myeloid/lymphoid or mixed-lineage leukem		7.60	24.00		
50	100761 100830	BE208491 AC004770	Hs.295112 Hs.4756	KIAA0618 gene product flap structure-specific endonuclease 1		7.00			7.99
	100867	U14622	FIS.4730	gb:Human transketolase-like protein gene		10.20			
	100902	M16029	Hs.287270	ret proto-oncogene (multiple endocrine n		8.00			
	100906	AU076916	Hs.5398	guanine monphosphate synthetase					5.16
55	100960	J00124	Hs.117729	keratin 14 (epidermolysis bullosa simple	2.57				
	101045	J05614		gb:Human proliferating cell nuclear anti					4.69
	101061	NM_000175	Hs.180532	glucose phosphate Isomerase					4.19
		L02840	Hs.84244	potassium voltage-gated channel, Shab-re	0.40	12.91			
60	101124	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	3.12				
60	101175	U82671	Hs.36980	melanoma antigen, family A, 2	3.50				5.69
	101181	BE262621 L24203	Hs.73798 Hs.82237	macrophage migration Inhibitory factor (ataxia-telangiectasia group D-associated	4.08				0.00
	101204 101210	L29301	Hs.2353	opioid receptor, mu 1	4.00		6.40		
	101216	AA284166	Hs.84113	cyclin-dependent kinase inhibitor 3 (CDK	2.53				
65	101228	AA333387	Hs.82916	chaperonin containing TCP1, subunit 6A (7.90
	101233	AL135173	Hs.878	sorbitol dehydrogenase					4.45
	101273	Z11933	Hs.182505	POU domain, class 3, transcription facto	8.50				
	101342	U52112	Hs.182018	interleukin-1 receptor-associated kinase					4.17
70	101346	AI738616	Hs.77348	hydroxyprostaglandin dehydrogenase 15-(N				21.89	
70	101369	NM_000892	Hs.1901	kallikrein B, plasma (Fletcher factor) 1	2.74			12.80	
	101396	BE267931	Hs.78996	proliferating cell nuclear antigen	3.24				7.90
	101431	BE185289 NM_000424	Hs.1076 Hs.195850	small proline-rich protein 18 (comifin) keratin 5 (epidermolysis bullosa simplex	8.31				1.50
	101448	AL035668	Hs.73853	bone morphogenetic protein 2	0.01			38.80	
75	101466	BE262660	Hs.170197	glutamic-oxaloacetic transaminase 2, mit					4.01
, ,	101484		Hs.20315	interferon-induced protein with tetratri				12.00	
	101502	M26958		gb:Human parathyroid hormone-related pro	10.50				
	101505	AA307680	Hs.75692	asparagine synthetase					4.46
00	101526		Hs.154721	aconitase 1, soluble	4.02				4.05
80	101535	X57152	Hs.99853	fibrillarin				0.00	4.65
	101577	M34353	Hs.1041	v-ros avian UR2 sarcoma virus oncogene h	E4.00			9.09	
	101649		Hs.1690	heparin-binding growth factor binding pr	54.00 5.50				
	101663		Hs.2178	H2B histone family, member Q H2A histone family, member A	5.59 7.00				
85	101664	AA436989 L24498	Hs.121017 Hs.80409	growth arrest and DNA-damage-inducible,	1.00	7.60			
05	101003		110,00700	States and series and series an assembly					

	W	O 02/086	443						PCT/US02/1247	76
	101695	M69136	Hs.135626	chymase 1, mast cell	4.79					
	101724	L11690	Hs.620	bullous pemphigold entigen 1 (230/240kD)	15.21					
	101748	NM_001944	Hs.1925	desmogleln 3 (pemphigus vulgaris antigen	55.50				440	
5	101759	M80244	Hs.184601	solute carrier family 7 (cationic amino myeloid cell nuclear differentiation ant				18.57	4.10	
,	101771 101804	NM_002432 M86699	Hs.153837 Hs.169840	TTK protein kinase	4.50			10.57		
	101809	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	140.00					
	101833	AU076442	Hs.117938	collagen, type XVII, alpha 1	2.56					
10	101842	M93221	Hs.75182	mannose receptor, C type 1				12.80	c 00	
10	101851	BE260964	Hs.82045	midkine (neurite growth-promoting factor		7.00			5.88	
	102002		Hs.81469 Hs.306098	nucleotide binding protein 1 (E.coli Min aldo-keto reductase family 1, member C1		7.80			4.35	
	102039 102072		Hs.78743	zinc finger protein 131 (clone pHZ-10)			7.40		4.00	
	102083		Hs.75117	interleukin enhancer binding factor 2, 4					5.12	
15	102111	L36198	Hs.81884	sulfotransferase family, cytosolic, 2A,				12.00		
	102123	NM_001809	Hs.1594	centromere protein A (17kD)	6.20					
	102154	U17760	Hs.75517	laminin, beta 3 (nicein (125kD), kalinin	2.62					
	102193 102217		Hs.313 Hs.301613	secreted phosphoprotein 1 (osteopontin, JTV1 gene	5.85				6.18	
20	102224	NM_002810	Hs.148495	proteasome (prosome, macropain) 26S subu					4.49	
	102234	AW163390	Hs.278554	heterochromatin-like protein 1					5.80	
	102251	NM_004398	Hs.41706	DEAD/H (Asp-Glu-Ala-Asp/His) box polypep	4.50					
	102305	AL043202	Hs.90073	chromosome segregation 1 (yeast homolog)					5.15	
25	102330 102340	BE298063	Hs.77254	chromobox homolog 1 (Drosophila HP1 beta macrophage stimulating 1 (hepatocyte gro				9.33	4.17	
23	102340	U37055 U37519	Hs.278657 Hs.87539	aldehyde dehydrogenase 3 family, member	8.87			3.00		
	102368	U39817	Hs.36820	Bloom syndrome	15.91					
	102394	NM_003816	Hs.2442	a disintegrin and metalloproteinase doma			19.20			
20	102404	NM_005429	Hs.79141	vascular endothelial growth factor C				14.00		
30	102537	U57094	Hs.50477	RAB27A, member RAS oncogene family				12.00	A 57	
	102581	AU077228 Al435128	Hs.77256 Hs.181369	enhancer of zeste (Drosophila) homolog 2 ubiquitin fusion degradation 1-like					4.57 3.98	
	102610	U65011	Hs.30743	preferentially expressed antigen in mela	77.50				0.00	
	102623	AW249285	Hs.37110	melanoma antigen, family A, 9	12.50					
35	102642	AA205847	Hs.23016	G protein-coupled receptor			22.00			
	102654	AV649989	Hs.24385	Human hbc647 mRNA sequence		12.00		40.00		
	.102659	BE245169	Hs.211610 Hs.29279	CUG triplet repeat, RNA-binding protein eyes absent (Drosophila) homolog 2	6.50			12.80		
	102669 102672	U71207 U72066	Hs.29279	retinoblastoma-binding protein 8	8.50				•	
40	102687	NM_007019	Hs.93002	ubiquitin carrier protein E2-C					9.24	
	102696	BE540274	Hs.239	forkhead box M1					5.54	
	102768	U82321		gb:Homo sapiens clone 14.9B mRNA sequenc		6.60			0.70	
	102781		Hs.108809	chaperonin containing TCP1, subunit 7 (e					3.78 4.26	
45	102784 102824	U85658 U90916	Hs.61796 Hs.82845	transcription factor AP-2 gamma (activat Homo sapiens cDNA: FLJ21930 fis, clone H			14.40		4.20	
13	102829	NM_006183	Hs.80962	neurotensin	8.00		14140		*	
	102888	Al346201	Hs.76118	ubiquitin carboxyl-terminal esterase L1					5.50	
		BE440042	Hs.83326	matrix metalloproteinase 3 (stromelysin			6.70			
50	102913	NM_002275	Hs.80342	keratin 15 small nuclear ribonucleoprotein polypept	4.64 2.93					
50	102935	. BE561850 X15218	Hs.80506 Hs.2969	v-ski avian sarcoma viral oncogene homol	2.53			11.40		
	102531	BE387202	Hs.118638	non-metastatic cells 1, protein (NM23A)				111.70	7.26	
	103023	AW500470	Hs.117950	multifunctional polypeptide similar to S	3.01					
F	103036	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial	27.90					
55 .	103038	AA926960	Hs.334883	CDC28 protein kinase 1					8.79 4.27	
	103060 103099	NM_005940 AI693251	Hs.155324 Hs.8248	matrix metalloproteinase 11 (stromelysin NADH dehydrogenase (ubiquinone) Fe-S pro		9.80			4.27	
	103119	X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta	4.05	0.00				
	103168	X53463	Hs.2704	glutathione peroxidase 2 (gastrointestin	3.07					
60	103185		Hs.74368	transmembrane protein (63kD), endoplasmi					5.62	
		M22440	Hs.170009	transforming growth factor, alpha		7.40	•		4.70	
	103223 103242	BE275607 ¥76342	Hs.1708 Hs.389	chaperonin containing TCP1, subunit 3 (g alcohol dehydrogenase 7 (class IV), mu o			100.00		4.70	
	103316		Hs.324728	SMA5			100.00	9.80		
65	103375		Hs.54416	sine oculis homeobox (Drosophila) homolo	9.71					
	103376		Hs.323378	coated vesicle membrane protein	14.00					
		NM_007069	Hs.37189	similar to rat HREV107	2.93			11.00		
	103391 103404	X94453 BE394784	Hs.114366 Hs.78596	pyrroline-5-carboxylate synthetase (glut proteasome (prosome, macropain) subunit,	2.93				5.15	
70	103430		Hs.20716	translocase of inner mitochondrial membr					3.98	
	103446	X98834	Hs.79971	sal (Drosophila)-like 2				21.40		
	103476		Hs.293007	aminopeptidase puromycin sensitive		13.00				
	103477		Hs.119018	transcription factor NRF	E 02		6.40			
75	103478 103515	BE514982 Y10275	Hs.38991 Hs.56407	S100 calcium-binding protein A2 phosphoserine phosphatase	5.02 10.50					
, 5	103515	BE616547	Hs.2785	keratin 17	6.41					
	103580	AA328046	Hs.46405	polymerase (RNA) II (DNA directed) polyp					3.84	
	103587	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein	78.50					
80	103594	Al368680	Hs.816	SRY (sex determining region Y)-box 2	6.51					
οv	103636 103768	NM_006235 AF086009	Hs.2407	POU domain, class 2, associating factor gb:Homo saptens full length insert cDNA	3.50				4.48	
	103766	AA314821	Hs.38178	hypothetical protein FLJ23468		8.00				
	103847	AF219946	Hs.102237	tubby super-family protein		10.40				
95	103913	AW967500	Hs.133543	ESTS			c ca	15.60		
85	104094	AA418187	Hs.330515	ESTs			6.60			

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	104150	AL122044	Hs.331633	hypothetical protein DKFZp566N034				26.00	1 C1/U302/124/0
	104257		Hs.9222	estrogen receptor binding site associate		6.80			3.98
	104261 104331	AW248364 AB040450	Hs.5409 Hs.279862	RNA polymerase I subunit cdk inhibitor p21 binding protein		6.80			3.80
5	104415	BE410992	Hs.258730	heme-regulated initiation factor 2-alpha		10.29			
	104558	R56678	Hs.88959	hypothetical protein MGC4816	4.21			15.79	
	104590 104658	AW373062 AA360954	Hs.83623 Hs.27268	nuclear receptor subfamily 1, group I, m Homo sapiens cDNA: FLJ21933 fis, clone H				17.40	
10	104660	BE298665	Hs.14846	Homo saplens mRNA; cDNA DKFZp564D016 (fr	6.40				
10	104689	AA420450	Hs.292911	ESTs, Highly similar to S60712 band-6-pr				10.00	6.55
	104754 104758	A1206234 BE560269	Hs.155924 Hs.7010	cAMP responsive element modulator NPD002 protein				10.00	4.47
	104971	BE311926	Hs.15830	hypothetical protein FLJ12691	2.87				
15	105011	BE091926	Hs.16244	mitotic spindle coiled-coil related prot	3.83				
13	105012 105026	AF098158 AA809485	Hs.9329 Hs.124219	chromosome 20 open reading frame 1 hypothetical protein FLJ12934	2.86	11.00			
	105076	A1598252	Hs.37810	hypothetical protein MGC14833					5.01
	105132	AA148164	Hs.247280	HBV associated factor			11.00		3.99
20	105143 105158	Al368836 AW976357	Hs.24808 Hs.234545	ESTs, Weakly similar to 138022 hypotheti hypothetical protein NUF2R		16.00	11.00		
_•	105175	AA305384	Hs.25740	ERO1 (S. cerevisiae)-like	4.32				,
	105200	AA328102	Hs.24641	cytoskeleton associated protein 2	3.00			40.00	
	105264 105298	AA227934 BE387790	Hs.26369	gb:zr57e08.s1 Soares_NhHMPu_S1 Homo sapi hypothetical protein FLJ20287	3.69			10.00	
25	105409	AW505076	Hs.301855	DiGeorge syndrome critical region gene 8				9.20	
	105460	AW296078	Hs.271721	Homo sapiens, clone IMAGE:4179986, mRNA,	4.40		7.80		
	105667 105743	AA767526 BE246502	Hs.22030 Hs.9598	paired box gene 5 (B-cell lineage specif sema domain, immunoglobulin domain (Ig),	4.12 3.82				
20	105782	H09748	Hs.57987	B-cell CLL/lymphoma 11B (zinc finger pro			27.00		
30	105848	AW954064	Hs.24951	ESTs			7.60		444
	105891 106019	U55984 AF221993	Hs.289088 Hs.46743	heat shock 90kD protein 1, alpha McKusick-Kaufman syndrome			16.80		4.14
	106069	BE566623	Hs.29899	ESTs, Weakly similar to G02075 transcrip			23.40		
25	106073	AL157441	Hs.17834	downstream neighbor of SON	9.50				
35	106126 106159	AA576953 AK001301	Hs.22972 Hs.3487	hypothetical protein FLJ13352 hypothetical protein FLJ10439	6.00				3.95
	106220	D61329	Hs.32196	mitochondrial ribosomal protein L36					6.04
	106260	AI097144	Hs.5250	ESTs, Weakly similar to ALU1_HUMAN ALU S			13.20		r.00
40	106300 106307	Y10043 AA436174	Hs.19114 Hs.37751	high-mobility group (nonhistone chromoso ESTs, Weakly similar to putative p150 (6.60			5.02
	106318	AA025610	Hs.9605	cleavage and polyadenylation specific fa		0.00			5.04
	106341	AF191020	Hs.5243	hypothetical protein, estradiol-induced	•		40.00		7.25
	106440 106481	AA449563 D61594	Hs.151393 Hs.17279	glutamate-cysteine ligase, catalytic sub tyrosylprotein sulfotransferase 1	4.75		13.80		
45	106586	AA243837	Hs.57787	ESTs				10.84	
	106605	AW772298	Hs.21103	Homo sapiens mRNA; cDNA DKFZp564B076 (fr	20.00			45.60	
* *	106654 106785	AW075485 Y15227	Hs.286049 Hs.20149	phosphoserine aminotransferase deleted in lymphocytic leukemia, 1	28.00 3.00				
50	106813	C05766	Hs.181022	CGI-07 protein			11.40		•
50	106895	AK001826	Hs.25245	hypothetical protein FLJ11269		e ee	6.00		•
	106913 106919	Al219346 AW043637	Hs.86178 Hs.21766	M-phase phosphoprotein 9 ESTs, Weakly similar to ALU5_HUMAN ALU S		6.56			4.27
	107054	A1076459	Hs.15978	KIAA1272 protein				34.80	
55	107059 107098	BE614410 Al823593	Hs.23044 Hs.27688	RAD51 (S. cerevisiae) homolog (E coli Re	4.71			24.80	
55	107098	AU076640	Hs.15243	ESTs nucleolar protein 1 (120kD)				24.00	7.05
	107129	AC004770	Hs.4756	flap structure-specific endonuclease 1	2.60		•		
	107198	AV657225 D20426	Hs.9846 Hs.41639	KIAA1040 protein		19.20 7.60			•
60	107203 107217	AL080235	Hs.35861	programmed cell death 2 DKFZP586E1621 protein	9.50	7.00			
	107284	NM_005629	Hs.187958	solute carrier family 6 (neurotransmitte	2.71				
	107318 107516	T74445 X57152	Hs.5957 Hs.99853	Homo sapiens clone 24416 mRNA sequence fibrillarin			8.71		4.33
	107529	BE515065	Hs.296585	nucleolar protein (KKE/D repeat)					4.00
65	107728	AA019551	Hs.294151	Homo sapiens, clone IMAGE:3603836, mRNA,		10.80			
	107851 107901	AA022953 L42612	Hs.61172 Hs.335952	EST keratin 6B	3.40		8.00		
	107922	BE153855	Hs.61460	lg superfamily receptor LNIR	2.88				
70	107932	AW392555	Hs.18878	hypothetical protein FLJ21620	7.50				
70	108015 108056	AW298357 AA043675	Hs.49927 Hs.62633	protein kinase NYD-SP15 ESTs				23.40 12.80	
	-108075	A1867370	Hs.139709	hypothetical protein FLJ12572				12.80	
	108187	BE245374	Hs.27842	hypothetical protein FLJ11210		7.00			
75	108296 108305	N31256 AA071391	Hs.161623	ESTs gb:zm61e06.r1 Stratagene fibroblast (937		6.60		11.80	
	108393	AA075211		gb:zm86a08.r1 Stratagene ovarian cancer				11.80	
	108480	AL133092	Hs.68055	hypothetical protein DKFZp434I0428				20.80	
	108554 108573	AA084948 AA086005		gb:zn13b09.s1 Stratagene hNT neuron (937 gb:zl84c04.s1 Stratagene colon (937204)		6.40		25.40	
80	108584	AA088326	Hs.120905	Homo saplens cDNA FLJ11448 fis, clone HE		9.60		20.70	
	108597	AK000292	Hs.278732	hypothetical protein FLJ20285	0.00			14.60	
	108695 108699	AB029000 AA121514	Hs.70823 Hs.70832	KIAA1077 protein ESTs	3.00			10.00	
0.5	108700	AA121518	Hs.193540	ESTs, Moderately similar to 2109260A B c			11.00	. 3.00	•
85	108780	AU076442	Hs.117938	collagen, type XVII, alpha 1	11.21				

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	108810	AW295647	Hs.71331	hypothetical protein MGC5350	8.50					
	108816	AA130884	Hs.270501	ESTs, Moderately similar to ALU2_HUMAN		7.40				
	108857	AK001468	Hs.62180	aniilin (Drosophila Scraps homolog), act	4.00					
5	108860	AA133334	Hs.129911 Hs.24341	ESTs	6.09 3.00					
J	108937 109010	AL050107 NM_007240	Hs.44229	transcriptional co-activator with PDZ-bi dual specificity phosphatase 12	2.69					
	109121	BE389387	Hs.49767	NADH dehydrogenase (ubiquinone) Fe-S pro	2.00				4.53	
	109166	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines	10.58					
10	109227	AA766998	Hs.85874	Human DNA sequence from clone RP11-16L21		9.00				
10	109415	U80736	Hs.110826	trinucleolide repeat containing 9		51.40		44.00		
	109418	A1866946	Hs.161707	ESTS			17.60	11.00		
	109454 109502	AA232255 AW967069	Hs.295232 Hs.211556	ESTs, Moderately similar to A46010 X-li hypothetical protein MGC5487			9.49			
	109543	AA564994	Hs.222851	ESTs ·		12.67	0.10			
15	109648	H17800	Hs.7154	ESTs				10.40		
	109680	AB037734	Hs.4993	KIAA1313 protein			33.20			
	109700	F09609		gb:HSC33H092 normalized infant brain cDN			44.00	16.00		
	109704	A1743880	Hs.12876	ESTs			11.00	12.60		
20	109792 109981	R49625 BE546208	Hs.26090	gb:yg61f03.s1 Soares infant brain 1NIB H hypothetical protein FLJ20272	4.00			12.00		
20	109998	AL042201	Hs.21273	transcription factor NYD-sp10	4.00	7.80				
	110039	H11938	Hs.21907	histone acetyltransferase		7.00				
	110156	AA581322	Hs.4213	hypothetical protein MGC16207					4.24	
25	110500	AA907723	Hs.36962	ESTs	4.50	0.00				
25	110551 110561	AW450381 AA379597	Hs.14529 Hs.5199	ESTs HSPC150 protein similar to ubiquitin-con	3.06	8.60				
	110854	BE612992	Hs.27931	hypothetical protein FLJ10607 similar to	3.00	6.80				
	110886		Hs.72249	three-PDZ containing protein similar to			8.80			
••	110916	BE178102	Hs.24349	ESTs		6.80				
30	111003	N52980	Hs.83765	dihydrofolate reductase				16.80		
	111337	AA837396	Hs.263925	LIS1-Interacting protein NUDE1, rat homo	2.54			0.00		٠,
	111434 111439		Hs.142736 Hs.19238	ESTs ESTs		•		9.80 10.40		
	111540		Hs.9786	ESTs zinc finger protein 275			15.40	10.40		
35	111597	R11499	Hs.189716	ESTs				9.20		
	111895	T80581	Hs.12723	Homo sapiens clone 25153 mRNA sequence		6.80				
	111929	AF027208	Hs.112360	prominin (mouse)-like 1				14.67		
	112054	R43590	11- 7004	gb:yc85g02.s1 Soares infant brain 1NIB H		10.80		40.20		
40	112210 112244	R49645 AB029000	Hs.7004 Hs.70823	ESTs KIAA1077 protein	2.99			10.20		
70	112382	R59904	115.70025	gb:yh07g12.s1 Soares infant brain 1NIB H	2.00	6.60				
	112392	R60763	Hs.193274	ESTs, Moderately similar to 157588 HSrei		5.55	7.10			
	112442		Hs.285681	Williams-Beuren syndrome chromosome regi	3.00					
15	112539	R70318	Hs.339730	ESTs				37.20		
45	112772	A1992283	Hs.35437	ESTs, Moderately similar to 138026 MLN 6				14.60	4.83	
	112869 112935	BE261750	Hs.4747 Hs.268760	dyskeratosis congenita 1, dyskerin ESTs	2.73				4.00	
		AA694010	Hs.6932	Homo sapiens clone 23809 mRNA sequence	2.70			12.00		
	112973	AB033023	Hs.318127	hypothetical protein FLJ10201	11.50					
50	112992	AL157425	Hs.133315	Homo sapiens mRNA; cDNA DKFZp761J1324 (f			10.89			
	113063	W15573	Hs.5027	ESTs, Weakly similar to A47582 B-cell gr	15.00		45.04			
	113073	N39342 T40444	Hs.103042	microtubule-associated protein 1B CAT56 protein		7.00	15.31			
	113078 113238	R45467	Hs.118354 Hs.189813	ESTs		7.00		41.20		•
55	113591	T91881	Hs.200597	KIAA0563 gene product				9.40		
	113702	T97307		gb:ye53h05.s1 Soares fetal liver spleen	25.00					
	113844	Al369275	Hs.243010	Homo sapiens cDNA FLJ14445 fis, clone HE			•	13.91		
	113984	R96696	Hs.35598	ESTS		7.80 7.20				
60	114073	R44953 AF155661	Hs.22908 Hs.22265	Homo sapiens mRNA; cDNA DKFZp434J1027 (f pyruvate dehydrogenase phosphatase	3.42	1.20				
00	114208	AL049466	Hs.7859	ESTs	0,72		6.74			
•	114251		Hs.21948	ESTs	•	-		33.20		
	114285,		Hs.22974	ESTs				13.20		
65	114313		Hs.27946	ESTs		7.00		10.00		
65	114339	AA782845 BE539976	Hs.22790	ESTS		7.80			4.14	
	114407 114560		Hs.103305 Hs.165221	Homo sapiens mRNA; cDNA DKFZp434B0425 (f ESTs				9.80	4,14	
		AA127386	110.100221	gb:zn90d09.r1 Stratagene lung carcinoma		7.60		0.00		
	114767	A1859865	Hs.154443	minichromosome maintenance deficient (S	3.21					
70	114793	AA158245		gb:zo76c03.s1 Stratagene pancreas (93720			6.00			
	114833		Hs.87159	hypothetical protein FLJ12577				11.40	4 24	
	115047	BE270930	Hs.82916 Hs.198249	chaperonin containing TCP1, subunit 6A (gap junction protein, beta 5 (connexin 3					4.31 4.03	
	115060 115097	AF052693 AA256213	Hs.72010	ESTs				35.40	4.00	
75	115113	AA256460		gb:zr81a04.s1 Soares_NhHMPu_S1 Homo sapi				15.20		
-	115123	AA256641	Hs.236894	ESTs, Highly similar to S02392 alpha-2-m					4.19	
		AW968073	Hs.194331	ESTs, Highly similar to A55713 inositol	05.00			12.40		
	115291	BE545072	Hs.122579	hypothetical protein FLJ10461	25.00	7.00				
80	115347	AA356792 AA662240	Hs.334824 Hs.283099	hypothetical protein FLJ14825 AF15q14 protein	3.25	7.00				
-	115522	BE614387	Hs.333893	c-Myc target JPO1	3.68					
	115536	AK001468	Hs.62180	anillin (Drosophila Scraps homolog), act	10.50					
	115566	A1142336	Hs.43977	Human DNA sequence from clone RP11-196N1				24.40		
85	115645	A1207410	Hs.69280	Homo sapiens, clone IMAGE:3636299, mRNA,	4.17		6.00			
O.J	115648	AW016811	Hs.234478	Homo sapiens cDNA: FLJ22648 fis, clone H			6.00			

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	115652	BE093589	Hs.38178	hypothetical protein FLJ23468	3.81	•				
	115697 115793	D31382 AA424883	Hs.63325 Hs.70333	transmembrane protease, serine 4 hypothetical protein MGC10753	62.14			11.80		
_	115816	BE042915	Hs.287588	Homo sapians cDNA FLJ13675 fis, clone PL				9.71		
5	115892		Hs.50831	ESTS	2.53		27.40			
	115906 115909	A1767756 AW872527	Hs.82302 Hs.59761	Homo sapiens cDNA FLJ14814 fis, clone NT ESTs, Weakly similar to DAP1_HUMAN DEATH	11.82					
	115965	AA001732	Hs.173233	hypothetical protein FLJ10970				34.29	0.22	
10	115978 115985	AL035864 AA447709	Hs.69517 Hs.268115	cDNA for differentially expressed CO16 g ESTs, Wealdy similar to T08599 probable	3.00				8.23	
10	116090	AJ591147	Hs.61232	ESTs	5.17					
	116096	AA682382 AF126743	Hs.59982	ESTs DNA I demain containing		10.60	8.20			
	116127 116157	BE439838	Hs.279884 Hs.44298	DNAJ domain-containing mitochondrial ribosomal protein S17		10.00			5.82	
15	116190	AI949095	Hs.67776	ESTs, Weakly similar to T22341 hypotheti	0.50				4.08	
	116278 116335	NM_003686 AK001100	Hs.47504 Hs.41690	exonuclease 1 desmocollin 3	9.50 3.67					
	116496	AW450694	Hs.21433	hypothetical protein DKFZp547J036	••••	7.00				
20	116503	AI925316 AI768015	Hs.212617 Hs.92127	ESTs ESTs			32.00	12.60		
20	116674 116929	AA586922	Hs.80475	polymerase (RNA) II (DNA directed) polyp		7.60	02.00			
	116973	Al702054	Hs.166982	phosphatidylinositol glycan, class F	•	9.80		10.20		
	116993 117079	Al417023 H92325	Hs.40478	ESTs gb:ys85f05.s1 Soares retina N2b4HR Homo				15.20		
25	117317	AI263517	Hs.43322	ESTs				13.40		
	117326 117396	N23629 W20128	Hs.241420 Hs.296039	Homo sapiens mRNA for KIAA1756 protein, ESTs				20.60 10.60		
	117412	N32536	Hs.42645	ESTs				16.00		
20	117519	N32528	Hs.146286	kinesin family member 13A				9.11	4.01	
30	117693 117721	AW179019 N46100	Hs.112110 Hs.93939	mitochondrial ribosomal protein L42 EST				19.80	4.01	
	117881	AF161470	Hs.260622	butyrate-induced transcript 1	2.71			47.00		
	117903 117992	AA768283 Al015709	Hs.47111 Hs.172089	ESTs Homo sapiens mRNA; cDNA DKFZp586l2022 (f				17.80	4.17	
35	118013	AI674126	Hs.94031	ESTs				10.60	7.7	
	118017	AI813444 N22886	Hs.42197	ESTs ESTs		7.00	8.82			
	118186 118325	AI868065	Hs.42380 Hs.166184	intersectin 2		7.00		13.80		
40	118367	N64269	Hs.48946	EST	244		6.14			
40	118368 118472	N64339 AL157545	Hs.48956 Hs.42179	gap junction protein, beta 6 (connexin 3 bromodomain and PHD finger containing, 3	3.14		12.40			
	118709	AA232970	Hs.293774	ESTs				12.20		•
	119025 119027	BE003760 AF086161	Hs.55209 Hs.114611	Homo sapiens mRNA; cDNA DKFZp434K0514 (f hypothetical protein FLJ11808	4.50 3.22					
45	119052	R10889	113.114011	gb:yf38d02.s1 Sozres fetal liver spleen	U.LL	9.60			•	•
	119164	AF221993	Hs.46743	McKusick-Kaufman syndrome			6.60	10.80		
	119186 119243	Al979147 T12603	Hs.101265	hypothetical protein FLJ22593 gb:CHR90123 Chromosome 9 exon il Homo sa				9.44		
50	119490	AA195276	Hs.263858	ESTs, Moderately similar to B34087 hypot		•	44.00	11.80		
50	119499 119599	Al918906 W45552	Hs.55080	ESTs gb:zc26d03.s1 Soares_senescent_fibroblas		12.60	14.80			
	119780	NM_016625	Hs.191381	hypothetical protein	17.00			,		
	119845 119941	W79123 AA699485	Hs.58561 Hs.58896	G protein-coupled receptor 87 ESTs	13.50	8.00				
55	119994	AA642402	Hs.59142	ESTs	7.73	0.00				
	120102	W67353	Hs.170218	KIAA0251 protein	2.91		39.60			
	120104 120294	AK000123 AK000059	Hs.180479 Hs.153881	hypothetical protein FLJ20116 Homo sapiens NY-REN-62 antigen mRNA, par	2.51		8.20			
60	120486	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog	8.73	7.00				
60	120599 120699	AA804448 Al683243	Hs.104463 Hs.97258	ESTs ESTs, Moderately similar to S29539 ribos		7.00		10.00	* :	·
	120715	AA292700		gb:zs59a06.s1 NCI_CGAP_GCB1 Homo sapiens		9.40			•	
	120821 120859	Y19062 AA826434	Hs.96870 Hs.1619	staufen (Drosophila, RNA-binding protein achaete-scute complex (Drosophila) homol		9.00		13.80		
65	120880	AA360240	Hs.97019	EST		15.60				
	120983		Hs.97587	EST			27.66 20.80			
	121034 121121	AL389951 AA399371	Hs.271623 Hs.189095	nucleoporin 50kD similar to SALL1 (sal (Drosophila)-like		22.80	20.00			
70	121313	AA402713	Hs.97872	ESTs	05.74			10.00		
70	121369 121376		Hs.128791 Hs.187958	CGI-09 protein solute carrier family 6 (neurotransmitte	25.71				5.42	
	- 121476	AA412311	Hs.97903	ESTs		8.30				
	121509 121553	AA868939 AA412488	Hs.97888 Hs.48820	ESTs TATA box binding protein (TBP)-associat	18.50	8.59				
75	121753	AK000552	Hs.323518	WD repeat domain 5	7.00					
	121838		Hs.98441	ESTS Lighby similar to A 35661 DNA excis	6.00			10.40		
	121857 121991		Hs.280858 Hs.98649	ESTs, Highly similar to A35661 DNA excis EST	0.00			12.20		
οΛ	122089	AW016543	Hs.98682	hypothetical protein FKSG32			8.60			
80	122105 122163		Hs.98699 Hs.98829	ESTs EST			6.14	10.40		
	122318	AA429743		gb:zv60b05.r1 Soares_testis_NHT Homo sap	40			18.20		
	122335 122338	AA443258 AA443311	Hs.241551 Hs.98998	chloride channel, calcium activated, fam ESTs	13.50 4.80					
85	122414		Hs.99087	ESTs, Weakly similar to S47073 finger pr		8.00		•		

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	122512	AF053305	Hs.98658	budding uninhibited by benzimidazoles 1			8.80		
	122516.	AA449352	Hs.99217	ESTs				9.40	
		A1220089	Hs.99439	EST ₆		9.20		10.40	
5	122852 122925	AI580056 AW268962	Hs.98992 Hs.111335	ESTs ESTs		6.80		10.40	
,	123005	AW369771	Hs.52620	integrin, beta 8		0.00	12.60		
	123044	AK001035	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro					5.35
	123160	AA488687	Hs.284235	ESTs, Weakly similar to 138022 hypotheti			6.06		
10	123315	AA495369	11- 470240	gb:zv37d10.s1 Soares ovary tumor NbHOT H			12.40 11.80		
10	123329 123497	Z47542 AA765256	Hs.179312 Hs.135191	small nuclear RNA activating complex, po ESTs, Weakly similar to unnamed protein		12.00	11.00		
	123518	AL035414	Hs.21068	hypothetical protein			13.00		
	123519	AW015887	Hs.112574	ESTs		12.20			
1.5	123614	AK000492	Hs.98806	hypothetical protein			7.80	10.60	
15	123616	AA680003	Hs.109363	Homo sapiens cDNA: FLJ23603 fis, clone L ESTs, Weakly similar to T2D3_HUMAN TRANS	23.00			10.00	
	123673 123727	BE550112 AI083986	Hs.158549 Hs.282977	hypothetical protein FLJ13490	25.00	7.00			
	123731	AA609839	110.202511	gb:ae62f01.s1 Stratagene lung carcinoma			9.80		
	123752	AA227714	Hs.179703	KIAA0129 gene product	3.50			40.00	
20	123900	AA621223	Hs.112953	EST	07.00			12.80	
	124006 124059	Al147155 BE387335	Hs.270016 Hs.283713	ESTs ESTs, Weakly similar to S64054 hypotheti	97.00 3.02				
	124069	AF134160	Hs.7327	claudin 1	0.02		27.80		
	124191	T96509	Hs.248549	ESTs, Moderately similar to S65657 alpha				35.80	
25	124273	AA457211	Hs.8858	bromodomain adjacent to zinc finger doma		7.20		44.00	
	124297	AL080215	Hs.102301	Homo sapiens mRNA; cDNA DKFZp586J0323 (f				11.00 16.00	
•	124305 124676	AW963221 Al360119.com	nHe 181013	gb:EST375294 MAGE resequences, MAGH Homo phosphoglycerate mutase 1 (brain)				10.00	6.08
	124874	BE550182	Hs.127826	RaiGEF-like protein 3, mouse homolog				21.00	
30	124904	AK000483	Hs.93872	KIAA1682 protein		9.40			•
	124969	Al650360	Hs.100256	ESTs				10:80	
	125000	T58615	Hs.110640	ESTs ESTs, Weakly similar to T33296 hypotheti		7.60		9.80	
	125201 125266	AA693960 W90022	Hs.103158 Hs.186809	ESTs, Highly similar to LCT2_HUMAN LEUKO		6.59			
35	125299	T32982	Hs.102720	ESTs				9.57	
	125356	AI057052	Hs.133554	ESTs, Weakly similar to Z195_HUMAN ZINC				14.00	
	125370	AA256743	Hs.134158	Homo saplens, Similar to KIAA0092 gene p			8.20	13.20	
	125418	AA777690 AL162066	Hs.188501 Hs.54320	ESTs hypothetical protein DKFZp762D096		21.40		13.20	
40	125433 125437	AL162000 Al609449	Hs.140197	ESTs		6.96			
	125446	BE219987	Hs.166982	phosphatidylinositol glycan, class F		8.80			
	125711	AA305800	Hs.5672	hypothetical protein AF140225				11.20	4.04
	125756	BE174587	Hs.289721	growth arrest specific transcript 5				15.60	4.31
45	125757 125769	A1274906 BE270266	Hs.166835 Hs.82128	ESTs, Highly similar to 1814460A p53-ass 5T4 oncofetal trophoblast glycoprotein	3.20			10.00	
73	125839	AW836261	Hs.337717	ESTs	0.20	8.20			
	125850	W85858	Hs.99804	ESTs	2.65				•
	125875	H14480		gb:ym18b09.r1 Soares infant brain 1NIB H		7.40			4.22
50	125924	BE272506	Hs.82109	syndecan 1					4.23 3.98
30	125972 126034	AI927475 H60340	Hs.35406	ESTs, Highly similar to unnamed protein gb:yr39b04.r1 Soares fetal liver spleen				10.60	
	126327	AA432266	Hs.44648	ESTs		11.60			
	126345	N49713		gb:yv23f06.s1 Soares fetal liver spleen		6.67		40.00	•
55	126435	AW614529	Hs.285847	CGI-19 protein				10.60	4.38
55	126487	AA283809	Hs.184601 Hs.203933	solute carrier family 7 (cationic amino ESTs		6.60			4.30
	126521 126522	Al475110 W31912	NS.200300	ob:zc76d03.s1 Pancreatic Islet Homo sapi		0.00		14.80	
	126543	AL035864	Hs.69517	cDNA for differentially expressed CO16 g					4.01
C O	126567	AA058394	Hs.57887	ESTs, Weakly similar to KIAA0758 protein			7.80	44.00	
60	126605	AA676910	11- 00007	gb:zj65h07.s1 Soares_fetal_liver_spleen_				11.60 14.60	
	126627 126628	AA497044 N49776	Hs.20887 Hs.170994	hypothetical protein FLJ10392 hypothetical protein MGC10946	8.00			14.00	
	126737	AW976516	Hs.283707	Homo sapiens cDNA: FLJ21354 fis, clone C	2.92				
	126795	AW975076	Hs.172589	nuclear phosphoprotein similar to S. cer	7.50				
65	126802	AW805510	Hs.97056	hypothetical protein FLJ21634		11.60			
	126892	AF121856	Hs.284291	sorting nexin 6	3.50			22.83	
	126928 126979	AA480902 AA210954	Hs.137401	ESTs gb:zq89h10.r1 Stratagene hNT neuron (937				11.80	
	126986	Al279892	Hs.46801	sorting nextn 14				11.60	
70	126992	Al809521		gb:wf30e03.x1 Soares_NFL_T_GBC_S1 Homo s				20.80	
	127066	R25066		gb:yg42c07.r1 Soares infant brain 1NIB H				27.60	
	127099	AA347668	U. 202505	gb:EST54026 Fetal heart II Homo sapiens				21.60 11.20	
•	127139 127209	AA830233 AA305023	Hs.293585 Hs.81964	ESTs SEC24 (S. cerevisiae) related gene famil	3.10			11120	
75	127221	BE062109	Hs.241551	chloride channel, calcium activated, fam	2.76				
-	127225	AA315933	Hs.120879	ESTs	40.00			16.80	
	127313	AK002014	Hs.47546	Homo sapiens cDNA FLJ11458 fis, clone HE	14.00			13.60	
	127444	AW978474 AW971353	Hs.7560 Hs.162115	Homo sapiens mRNA for KIAA1729 protein, ESTs		11.20		10.00	
80	127500 127524	A1243596	Hs.94830	ESTs. Moderately similar to T03094 A-kin			7.80		
	127540	N45572	Hs.105362	Homo sapiens, clone MGC:18257, mRNA, com	3.53				
	127599	AA613204	Hs.150399	ESTs				13.80	
	127609	X80031	Hs.530	collagen, type IV, atpha 3 (Goodpasture				28.00 19.80	
85	127662 127668	W80755 Al343257	Hs.8294 Hs.139993	KIAA0196 gene product ESTs				11.20	
55	121000	THETOLUI	103333						

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	127746		Hs.120189	ESTs				14.18	
	127812	AA741368	Hs.291434	ESTs .	4.50				
	127817	AA836641	Hs.163085	ESTS				24.60 9.20	
5	127959 127960		Hs.124292 Hs.41569	Homo sapiens cDNA: FLJ23123 fis, clone L phosphatidic acid phosphatase type 2A				16.83	
•	127969	F06498	Hs.93748	Homo sapiens cDNA FLJ14676 fis, clone NT		13.60			
	128015		Hs.334659	hypothetical protein MGC14139		7.00		DT 40	
	128027	AJ433721	Hs.164153	ESTs				37.40 9.60	
10	128077 128168	AI310330 NM_006147	Hs.128720 Hs.11801	ESTs Interferon regulatory factor 6				9.24	
10	128226	AI284940	Hs.289082	GM2 ganglloside activator protein	19.00				
		A1954968	Hs.279009	matrix Gla protein				10.40	
	128341		Hs.185030	ESTs		9.00			4.30
15	128527 128539	AA504583 R46163	Hs.101047 Hs.258618	transcription factor 3 (E2A immunoglobul ESTs		12.60		,	4.50
13	128568		Hs.274691	adenylate kinase 3		12.00			4.56
		AA933022	Hs.256583	interleukin enhancer binding factor 3, 9				10.00	
	128777	AI878918	Hs.10526	cysteine and glycine-rich protein 2			16.80		4.40
20	128781		Hs.105465 Hs.105924	small nuclear ribonucleoprotein polypept defensin, beta 2		8.12			4.48
20	128796 128920		Hs.166468	programmed cell death 5		0.12			4.62
	128924		Hs.26557	plakophilin 3					4.04
	128971	H05132	Hs.107510	ESTs		12.60			
-25		AL079648	Hs.301088	ESTs solute carrier family 2 (facilitated glu		8.80		•	6.05
23	129041	BE382756 BE250162	Hs.169902 Hs.83765	dihydrofolate reductase	2.59				0.00
		AI769160	Hs.108681	Homo sapiens brain tumor associated prot	•		6.67		
		AB023179	Hs.9059	KIAA0962 protein		8.00			
30		AF013758	Hs.109643	polyadenylate binding protein-interactin	4.00				4.06
30	129241 129300		Hs.109706 Hs.110165	hematological and neurological expressed ribosomal protein L26 homolog	2.55				4.00
	129404		Hs.317584	ESTs	18.00				
	129457	X61959	Hs.207776	aspartylglucosaminidase	6.50				
25	129466		Hs.334309	keratin 6A	12.94 -			11.00	
35		AJ148976 AF061812	Hs.112062 Hs.115947	ESTs keratin 16 (focal non-epidermolytic palm				11.00	4.46
	129641		Hs.11805	ESTs				12.00	
		AW163331	Hs.118778	KDEL (Lys-Asp-Glu-Leu) endoplasmic retic					4.70
40	129703		Hs.179999	Homo saplens, clone IMAGE:3457003, mRNA					4.02 5.71
40	129720 129748	AA156214 M16707	Hs.12152 Hs.123053	APMCF1 protein H4 histone, family 2	3.50				5.71
	129890		Hs.282804	hypothetical protein FLJ22704	0.00				4.21
	129896	BE295568	Hs.13225	UDP-Gal:betaGlcNAc beta 1,4- galactosylt	2.56				
15	129945	BE514376	Hs.165998	PAI-1 mRNA-binding protein			7.00		4.03
45	130010 130026	AA301116	Hs.142838 Hs.332112	nucleolar phosphoprotein Nopp34 EST		6.40	7.00		•
	130020		Hs.147097	H2A histone family, member X		0.40			4.65
	130149	AW067805	Hs.172665	methylenetetrahydrofolate dehydrogenase	2.74				
50		AA063546	Hs.75981	ubiquitin specific protease 14 (tRNA-gua			7.40		3.91
50	130441	U63630 AW409701	Hs.155637 Hs.1578	protein kinase, DNA-activated, catalytic bacutoviral IAP repeat-containing 5 (sur	4.87				3.91
	130500	AB007913	Hs.158291	KIAA0444 protein				9.60	
	130524	U89995	Hs.159234	forkhead box E1 (thyroid transcription f			13.40		
55	130541	X05608	Hs.211584	neurofilament, light polypeptide (68kD)			8.20		6.06
22	130553 130567	AF062649 AA383092	Hs.252587 Hs.1608	pituitary tumor-transforming 1 replication protein A3 (14kD)			7.00		0.00
	130577	M69241	Hs.162	insulin-like growth factor binding prote	3.04				
	130627	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage	3.87				
6 0	130648	AI458165	Hs.17296	hypothetical protein MGC2376				16.20 17.80	
60	130697 130744		Hs.1802 Hs.18747	major histocompatibility complex, class POP7 (processing of precursor, S. cerevi				17.00	5.28
	130800		Hs.19574	hypothetical protein MGC5469					4.43
	130867	NM_001072	Hs.284239	UDP glycosyltransferase 1 family, polype	16.84				
65	130869	J03626	Hs.2057	uridine monophosphale synthetase (orotat				9.60	4.92
65	130925 130994		Hs.169378 Hs.327337	multiple PDZ domain protein ESTs		12.40		3.00	
	131028	AI879165	Hs.2227	CCAAT/enhancer binding protein (C/EBP),	10.21				
	131031	NM_001650	Hs.288650	aquaporin 4				9.80	,
70	131041		Hs.22452	Homo sapiens mRNA for KIAA1737 protein,				9.60	·
70	131058	W28545 Al143139	Hs.101514 Hs.2288	hypothelical protein FLJ10342 visinin-like 1	2.74			17.00	
		H15302	Hs.168950	Homo sapiens mRNA; cDNA DKFZp566A1046 (f			8.80		
		AW953575	Hş.303125	p53-induced protein PIGPC1	3.12				
75		BE280074	Hs.23960	cyclin B1	3.07				
13	131200	BE540516 W25005	Hs.293732 Hs.24395	hypothetical protein MGC3195 small inducible cytokine subfamily B (Cy	3.07 2.87				
	131219		Hs.24908	ESTs				14.67	
	131375	AW293165	Hs.143134	ESTs			19.20		
80	131460		Hs.27076	RNA 3'-terminal phosphate cyclase	3.50				
οU	131476 131510		Hs.334644 Hs.27842	hypothetical protein FLJ14668 hypothetical protein FLJ11210	15.00		7.80		
	131646		Hs.30057	MRS2 (S. cerevisiae)-like, magnesium hom			7.00		
	131786	BE000971	Hs.306083	Novel human gene mapping to chomosome 22	2.65				
85	131839		Hs.33010	KIAA0633 protein				35.20	4.11
0,5	131043	AA192315	Hs.184062	putative Rab5-interacting protein					***

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	131877	O 02/086 J04088	443 Hs.156346	topoisomerase (DNA) II alpha (170kD)	19.00				PC1/USU2/124/0
	131885	BE502341	Hs.3402	ESTs	6.48				
	131921		Hs.34720	ESTs	EC 00		8.40		
5	131945 131958	NM_002916 NM_014062	Hs.35120 Hs.3566	replication factor C (activator 1) 4 (37 ART-4 protein	56.00				3.82
_	131965	W79283	Hs.35962	ESTs	3.03				
	132000 132040	AW247017	Hs.36978	melanoma antigen, family A, 3 Homo sapiens cDNA: FLJ22373 fis, clone H	3.30	9.80			
	132109	NM_001196 AW190902	Hs.315689 Hs.40098	cysteine knot superfamily 1, BMP antagon	21.00				
10	132114	NM_006152	Hs.40202	lymphoid-restricted membrane protein		8.40	•		
	132162 132164	AA315805 Al752235	Hs.94560 Hs.41270	desmoglein 2 procollagen-lysine, 2-oxoglutarate 5-dio	2.70				12.25
	132180	NM_004460	Hs.418	fibroblast activation protein, alpha	2.71				
1.5	132181	AW961231	Hs.16773	Homo sapiens clone TCCCIA00427 mRNA sequ	3.83			40.00	
15	132182 132231	NM_014210 AA662910	Hs.70499 Hs.42635	ecotropic viral integration site 2A hypothetical protein DKFZp434K2435	9.50			13.20	
	132277	AK001745	Hs.184628	hypothetical protein FLJ 10883	4.50				
	132328	NM_014787	Hs.44896	DnaJ (Hsp40) homolog, subfamily B, membe				9.20	
20	132394 132424	AK001680 AA417878 -	Hs.30488 Hs.48401	DKFZP434F091 protein ESTs, Moderately similar to ALU8_HUMAN A			8.60	19.80	
20	132528	T78736	Hs.50758	SMC4 (structural maintenance of chromoso			27.40		•
	132543	BE568452	Hs.5101	protein regulator of cytokinesis 1	4.38	7.00			
	132544 132550	L19778 AW969253	Hs.51011 Hs.170195	H2A histone family, member P bone morphogenetic protein 7 (osteogenic	2.64	7.00			•
25	132552	BE621985	Hs.296922	thiopurine S-methyltransferase				15.83	
	132581	AK000631	Hs.52256	hypothetical protein FLJ20624	4.00		6.60		
	132617 132638	AF037335 AJ796870	Hs.5338 Hs.54277	carbonic anhydrase XII DNA segment on chromosome X (unique) 992	4.95	8.20			
20	132653	Z15008	Hs.54451	laminin, gamma 2 (nicein (100kD), kalini	4.38				
30	132669	W38586	Hs.293981	guanine nucleotide binding protein (G pr	4.00				4.36
	132710 132771	W74001 Y10275	Hs.55279 Hs.56407	serine (or cysteine) proteinase inhibito phosphoserine phosphatase	4.60 3.71				
	132799	W73311	Hs.169407	SAC2 (suppressor of actin mutations 2,	• • • • • • • • • • • • • • • • • • • •			9.48	
35	132833	U78525	Hs.57783	eukaryotic translation initiation factor				12.00	5.83
33	132892 132906	AW834050 BE613337	Hs.9973 Hs.234896	tensin geminin	3.09			12.00	
	132959	AW014195	Hs.61472	ESTs, Weakly similar to YAE6_YEAST HYPOT					3.87
	132962		Hs.6153	CGI-48 protein	3.50 6.18				•
40	132990 132994	X77343 AA112748	Hs.334334 Hs.279905	transcription factor AP-2 alpha (activat clone HQ0310 PRO0310p1	3.19				•
	133000	AL042444	Hs.62402	p21/Cdc42/Rac1-activated kinase 1 (yeast	2.96				
	133050 133083	X73424 BE244588	Hs.63788 Hs.6456	propionyl Coenzyme A carboxylase, beta p chaperonin containing TCP1, subunit 2 (b	2.55				4.00
	133086	L17131	Hs.139800	high-mobility group (nonhistone chromoso					8.96
45	133134	AF198620	Hs.65648	RNA binding motif protein 8A				40.00	4.28
	133155 133181	M58583 X91662	Hs.662 Hs.66744	cerebellin 1 precursor twist (Drosophila) homolog (acrocephalos	3.00			10.80	
	133204	BE267696	Hs.254105	enolase 1, (alpha)	0.00				4.63
50	133412		Hs.73112	guanine nucleotide binding protein (G pr	0.00	12.50			
30	133421 133451	AF134160 AW970026	Hs.7327 Hs.73818	claudin 1 ubiquinol-cytochrome c reductase hinge p	2.85				4.66
	133453	A1659306	Hs.73826	protein tyrosine phosphatase, non-recept		6.80			•
	133504	NM_004415	Hs.74316	desmoplakin (DPI, DPII)	6.14	•			4.55
55	133506 133615	BE562958 M62843	Hs.74346 Hs.75236	hypothetical protein MGC14353 ELAV (embryonic lethal, abnormal vision,				17.80	4.55
	133627	NM_002047	Hs.75280	glycyl-tRNA synthetase					4.85
	133649 133669	U25849 NM_006925	Hs.75393 Hs.166975	acid phosphatase 1, soluble splicing factor, arginine/serine-rich 5				14.00	6.34
	133749	L20852	Hs.100373	solute carrier family 20 (phosphate tran			6.11	17.00	
60	133776	BE268649	Hs.177766	ADP-ribosyltransferase (NAD+; poly (ADP-	0.0=				4.91
	133865 133946	AB011155 AJ001258	Hs.170290 Hs.173878	discs, large (Drosophila) homolog 5 NIPSNAP, C. elegans, homolog 1	. 3.07				4.60
	133973		Hs.78026	ESTs, Weakly similar to similar to ankyr				13.00	
65	134047	BE262529	Hs.78771	phosphoglycerate kinase 1	0.50				3.85
03	134098 134107	BE513171 NM_005629	Hs.79086 Hs.187958	mitochondrial ribosomal protein L3 solute carrier family 6 (neurotransmitte	2.56		8.20		
	134112	AW449809	Hs.79150	chaperonin containing TCP1, subunit 4 (d					4.08
	134158	U15174	Hs.79428	BCi_2/adenovirus E1B 19kD-interacting pro	31.00		24.60		
70	134160 134168	T98152 AA398908	Hs.79432 Hs.181634	fibrillin 2 (congenital contractural ara Homo sapiens cDNA: FLJ23602 fis, clone L			24.00		6.71
	134185	AA285136	Hs.301914	neuronal specific transcription factor D				14.74	,
	134201 134272	L35035 X76040	Hs.79886 Hs.278614	ribose 5-phosphate isomerase A (ribose 5 protease, serine, 15	4.50	8.40			
	134272	BE083936	Hs.80976	antigen identified by monoclonal antibod	******	9.00			
75	134353	AL138201	Hs.82120	nuclear receptor subfamily 4, group A, m	0.00			16.40	
	134367 134380	AA339449 AU077143	Hs.82285 Hs.179565	phosphoribosylglycinamide formyltransfer minichromosome maintenance deficient (S.	2.80 4.68				
	134423	H53497	Hs.83006	CGI-139 protein					3.84
80	134469	AA279661	Hs.83753	small nuclear ribonucleoprotein polypept					5.81
ου	134470 134498	X54942 AW246273	Hs.83758 Hs.84131	CDC28 protein kinase 2 threonyl-tRNA synthetase					4.21 7.30
	134502	BE148534	Hs.84168	UV-B repressed sequence, HUR 7		13.60			
	134510 134548	NM_002757 N95406	Hs.250870	mitogen-activated protein kinase kinase Deleted in split-hand/split-foot 1 regio				9.70	4.63
85	134548	N95406 AK001741	Hs.333495 Hs.8739	hypothetical protein FLJ 10879	6.00				
			·	••					

	W	O 02/086	443					PCT/US02/12476		
	134724	AF045239	Hs.321576	ring finger protein 22				12.00		
	134743	AA044163	Hs.89463	potassium large conductance calcium-acti	4.00					
	134781	AA374372	Hs.89626	parathyroid hormone-like hormone			25.20			
_	134806	AD001528	Hs.89718	spermine synthase					4.58	
5	134853	BE268326	Hs.90280	5-aminoimidazole-4-carboxamide ribonucle					4.79	
	134859	D26488	Hs.90315	KIAA0007 protein			6.20			
	134891	R51083	Hs.90787	ESTs			7.40			
	134960	BE246400	Hs.285176	acetyl-Coenzyme A transporter	4.00					
10	134993	BE409809	Hs.301005	purine-rich element binding protein B					4.48	
10	135047	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	9.50					
	135080	Al761180	Hs.94211	rcd1 (required for cell differentiation,	5.00					
	135103	NM_003428	Hs.9450	zinc finger protein 84 (HPF2)		11.00				
	135145	AW014729	Hs.95262	nuclear factor related to kappa 8 bindin					4.01	
1.5	135184	U13222	Hs.96028	forkhead box D1			7.00			
15	135242	Al583187	Hs.9700	cyclin E1	13.50					
	135286	AW023482	Hs.97849	ESTs	6.46					
	135289	AW372569	Hs.9788	hypothetical protein MGC10924 similar to		8.80				
	135355	AK001652	Hs.99423	ATP-dependent RNA helicase	10.00					
20	135371	NM_006025	Hs.997	protease, serine, 22	8.00			44.00		
20	135393	L11244	Hs.99886	complement component 4-binding protein,				14.60		

TABLE 5B shows the accession numbers for those primekeys lacking unigenelD's for Table 5A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

25	similarity using Cluster "Accession" column.	ring and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the
30	CAT number: Gene cl	Eos probeset identifier number uster number k accession numbers
	Pkey CAT num	ber Accessions
35	117079 1621717_ 124305 242163_1 101502 18202_6 109792 754958_1	AW963221 AA344870 AA344871 H93331 M26958 R49625 F10674
40	126034 1598157_ 102768 44641_1 126345 1653833_ 127066 1703458_ 127099 244301_1	U82321 H66077 1 N49713 N49819 W03810 1 R25066 R20144 R20145 Z43845 AA347668 AW956810 Z44271 F07065 F07064 R13506
45	119243 1774795_ 125875 1566433_ 112054 1538292_ 126979 171411_1 126992 880655_1	1 H14480 N98295 1 R43590 F10439 AA210954 AA211007
50	122318 292419_1 114699 135322_1 114793 150742_1 108305 111550_1 108393 113411_1	AA127386 R15644 AA127404 AA158245 AA158235 AA071391 AA069892 AA069891 AA075211 AA075245 AA075126 AA074946
55	109700 genbank 120715 genbank 113702 genbank	AA609839 AA609839 F09609 F09609 AA292700 AA292700 T97307 T97307
60	101045 entrez_J0 108554 genbank_	5614 J05614 AA084948 AA084948 AA086005 AA086005 R10889 R10888
65	126605 439280_1 103768 46922_1	AA676910 AA778853 AA778865 W86800 W42667 AI580740 AI690440 AI561350 AW467906 AW151450 AI825927 AL041716 AI885600 AI742213 AW248624 AI955498 AA033947 AA845593 AI623711 N68583 C00064 AA193567 AW083868 AW163216 AA191595 AA522778 AI628008 AI915518 AA843508 AI926195 AA176265 AW167963 AA992115 W93647 AW103572 AI862994 AI342059 AA911719 AA176155 AA024712 AA069988 AA205591 AI591107
70		A1199673 AIB11766 A1275832 A1422233 A1191852 A1096682 A1580124 A1683612 AA582453 AA927559 AA486415 T32414 A1084978 H44849 H44848 H20477 T91695 W47039 AA070055 AA024795 AA328855 AA379248 AA379330 AA385580 W25920 W03688 AA448359 AA093881 AW362477 AA089997 A1350265 W93479 N99688 AA932257 AW351469 H68590 AA663402 AA069771 AW087986 A1858420 AA6002144 A1970774 A1857712 A1683081 A1885584 AW131150 A1567981 AW002714 AW189973 AW075495 AW168303 AA953714 AW516881 A1357375 A1566663 AW512676 A1570580 A1023690 AA448216 A1079853 A1422707 AA779516 AW026972 AW130082 AW162307 AW438846 AA709332 AW192394 A1167350 A1217879 A1129152 AA719509 A1350480 AA663418 A1003634 AW118546 AA180261 AA442833 A1268625 AA888881
75		AI038759 AA846723 AI248770 AA993694 AI280335 AI885107 AW518649 AA641563 AA995835 AA582521 AI276744 AA436478 AI017360 AI620763 AI859887 N73926 AI076327 AI741615 AI160617 AW172819 AI492005 AA677429 AA996334 AI693771 AI950039 AI245629 AI288515 AI866186 T93293 AA173262 AA599779 AI680092 AW439316 AI084555 AI272672 AI583507 AW473219 AA738132 AW473283 AI367492 AA995410 AI689624 AA206353 AI033095 AI040382 AA873630 AI221074 AI934840 AI418680 AA844306 R94503 AA773520 AA843169 AA219425 AA629658 AI811719 AW411275 AI590981 W37907 AI591178 AI684051 AA983238 AA669347 AA976239 AA704570 AI628339
80		A1884391 A1241580 A1003539 AW176687 AA009650 N34566 A1333493 A1186070 AA070827 AA411683 A1280884 AA872023 AA207255 AA021576 N71953 A1885888 AW076039 T15777 A1537673 AW248048 H09554 W93480 W47001 AW079114 AA063160 AA57453 R60788 A1859431 H20478 AA218882 AA757465 AA100995 A1864135 A1934209 AA070503 H47008 AA219646 W61039 W93907 AW385050 W37967 W78028 AA189007 AA479136 R93650 AA442312 T30287 AA847628 AA180262 AA009549 C03892 AW149464 AA310963 AA219693 AA069747 R29207 AA094784 AA293615 AA447848 A1984167 N90393 C05097 N56499 AW292351 AW149681 AW473258 AA629322 A1004409
85		AW105577 AI954937 AI811070 AA902422 AW514437 AA535460 AA916877 AW517122 AA974657 AA975649 AW517130 AW517129 F31737 W07688 AA193645 AA378994 AA489273 F32267 W39303 AA021181 N86810 AA406524 AA062553 AA436801 H08985 H15979 N40310

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AA436789 AA232172 AW360778 W25862 R60282 AA436530 AA378894 AA187461 AI940535 AA604210 AA089514 AA360421 N88243 N84281

AA209340 N56174 N88374 AA191088 AW247691 AA249013 AA093111 AA972536 AW298594 AA375893 T12139 W28186 AW243849

AI288629 AA843996 W15260 AI188286 AW248079 R15836

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genbank_W45552 W45552
genbank_R59904
genbank_AA227934
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AA496369
AA496646 119599 5 112382 105264 100071 123315

	120010		,,,,,,,,	300 / 0 11300 10				
10	Eos/Affyrr	retrix Hu03 Gen	echip array. (nonsmokers with lung cancer relative to smokers with lung Gene expression data for each probeset obtained from this	g cancer. These gen analysis was expres	es were selected from t sed as average intensit	59680 probesets o y (AI), a normalize	on the ed value reflecti
	the relativ	e level of mRN/	A expression.					
15	Pkey: ExAccn: Unigenell	Exemplar	Accession nu	entifier number Imber, Genbank accession number				
	Uninone	Title: Unigene						
	R1:	average	yene uue nf Al for eamol	es from non-smokers with adenocarcinoma divided by the	90th nercentile of Al	for samples from smoke	ars with adenocan	cinoma
20	R2:	average of carcinom.	of Al for sampl	es from non-smokers with squamous cell carcinoma divide	d by the 90th percen	tile of AI for samples fro	ım smokers with s	quamous cell
	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2	-	
25	100971	BE379727	Hs.83213	fatty acid binding protein 4, adipocyte		3,64		
	101174	L17330	Hs.280	pre-T/NK cell associated protein	15.00			
	101296	Y12490	Hs.85092	thyroid hormone receptor interactor 11		2.46		
	101304	AA001021	Hs.6685	thyroid hormone receptor interactor 8		12.00		
	101806	AA586894	Hs.112408	S100 calcium-binding protein A7 (psorias		2.68		
30	101972	S82472		gb:beta -pol=DNA polymerase beta (exon a		2.11		
	102274	U30930	Hs.158540	UDP glycosyltransferase 8 (UDP-galactose	7.50			
	102394	NM_003816	Hs.2442	a disintegrin and metalloproteinase doma	7.50			
	102832	U92015		gb:Human clone 143789 defective mariner	13.50			
	103010	X52509	Hs.161640	tyrosine aminotransferase	9.50			
35	103439	X98266		gb:H.sapiens mRNA for ligase like protei		2.50		
	103563	L02911	Hs.150402	activin A receptor, type I	9.00			
	103857	Al076795 ·	Hs.45033	lacrimal proline rich protein		3.94		
	104239	AB002367	Hs.21355	doublecortin and CaM kinase-like 1	13.50	40.00		
40	104590	AW373062	Hs.83623	nuclear receptor subfamily 1, group I, m	40.00	12.66		
40	104907	AA055829	Hs.196701	ESTs, Weakly similar to ALU1_HUMAN ALU	16.50	0.47		
	106131	BE514788	Hs.296244	SNARE protein	7.00	2.17		
	106672	H47233	Hs.30643	ESTs	7.00			
	106872	T56887	Hs.18282	KIAA1134 protein	11.50	2.20		
15	106960	AA156238	Hs.32501	ESTs	0.50	2.38		*
45	106971	Z43846	Hs.194478	Homo sapiens mRNA; cDNA DKFZp43401572 (f	9.50	2.95		
	107982	AA035375	Hs.57887	ESTs, Weakly similar to KIAA0758 protei	16.50	2.50		
	108562	AA100796	11- 00220	gb:zm26c06.s1 Stratagene pancreas (93720	13.00			
	108599	AB018549	Hs.69328	MD-2 protein	13.00	2.40		
50	108663	BE219231	Hs.292653	ESTs, Wealty similar to T26845 hypotheti	7.00	2.40		
20	109247	AA314907	Hs.85950	ESTs ESTs	7.00	5.00		
	109630	R44607	Hs.22672	Homo sapiens mRNA; cDNA DKFZp434M082 (fr	12.50	0.00		
	110193	A1004874 H24458	Hs.310764 Hs.32085	EST	16.50			
	110234 110644	R94207	Hs.268989	ESTs, Highly similar to type II CALM/AF1	8.00			
55	110886	AW274992	Hs.72249	three-PDZ containing protein similar to	17.00			
55	111057	T79639	Hs.14629	ESTs	16.50			
	111950	AF071594	Hs.110457	Wolf-Hirschhorn syndrome candidate 1	11.00			
	112291	R53972	Hs.26026	ESTs		3.00		
	112956	Z43784	Hs.75893	ankyrin 3, node of Ranvier (ankyrin G)	•	2.79		:
60	113009	T23699	Hs.7246	ESTs		4.50		
•	113060	BE564162	Hs.250820	hypothetical protein FLJ14827	9.79		•	
	113073	N39342	Hs.103042	microtubule-associated protein 18	32.50		•	
	113074	AK001335	Hs.31137	protein tyrosine phosphatase, receptor t		3.82		
	113121	T48011	Hs.8764	EST		2.21		
65	113125	AA968672	Hs.8929	hypothetical protein FLJ11362	19.50			
	113757	AA703095	Hs.18631	ESTs		2.65		
	113848	W52854	Hs.27099	hypothetical protein FLJ23293 similar to	6.00			
	113884	A1333076	Hs.28529	chromosome 12 open reading frame 2		6.00		
	113936	W17056	Hs.83623	nuclear receptor subfamily 1, group I, m		4.63		
70	114875	AA235609	Hs.236443	Homo sapiens mRNA; cDNA DKFZp564N1063 (7.00		
	114987	AA251016	Hs.87808	EST		6.00		
	115460	AW958439	Hs.38613	ESTs		2.27		
	115722	W91892	Hs.59609	ESTs		9.00		
75	116261	AA481788	Hs.190150	ESTs	9.50			
75	116830	H61037	Hs.70404	ESTs, Weakly similar to ALU2_HUMAN ALU	8.50			
	116970	AB023179	Hs.9059	KIAA0962 protein	7.50	2 69		
	117178	H98675	Hs.269034	ESTs	7 50	2.68		
	117757	AF088019	Hs.46732	EST	7.50			
0Λ	118283	AA287747	Hs.173012	ESTs, Weakly similar to A46010 X-linked	16.50	2.50		
80	118384	AF217525	Hs.49002	Down syndrome cell adhesion molecule		2.50		
	118657	A1822106	Hs.49902	ESTs		2.39		
	120328	AA923278	Hs.290905	ESTs, Weakly similar to protease [H.sapi	7 00	3.50		
	120404	AB023230	Hs.96427	KIAA1013 protein	7.00 6.00			
85	120524	AA261852	Hs.192905	ESTs Homo sapiens cDNA: FLJ23004 fis, clone L	17.92			
05	120688	AW207555	Hs.97093	Fromito papierio contra i integorori ilo, ciona E	11.32			
					•			

	W	O 02/086	443				PCT/US02/12476
	121558	AA412497		gb:zt95g12.s1 Soares_testis_NHT Homo sap		2.95	
	121676	H56037	Hs.108146	ESTs	10.00		
	121936	AI024600	Hs.98612	ESTs	15.00		
	121938	AA428659	Hs.98610	ESTs	14.00		
5	122177	AA435789	Hs.98833	EST	8.93		
•	123442	AA299652	Hs.111496	Homo sapiens cDNA FLJ11643 fis, done HE	13.04		
	123551	AA608837	1101111100	ob:af03h12.s1 Soares_testis_NHT Homo sap	11.50		
	123756	AA609971	Hs.112795	EST	11.00		
	123861	AA620840		gb:af89g01.s1 Soares_testis_NHT Homo sap		2.50	
10	124371	N24924	Hs.188601	ESTs	6.50		
	127477	BE328720	Hs.280651	ESTs		4.33	
	127591	Al190540	Hs.131092	ESTs		3.02	
	128252	AA455924	Hs.192228	ESTs	7.00		
	128426	AI265784	Hs.145197	ESTs		2.08	
15	128925	R67419	Hs.21851	Homo sapiens cDNA FLJ12900 fis, clone NT		2.11	
	128945	AI990506	Hs.8077	Homo sapiens mRNA; cDNA DKFZp547E184 (fr	10.00		
	129105	Al769160	Hs.108681	Homo sapiens brain tumor associated prot	15.50		
	129235	AW977238	Hs.126084	KIAA1055 protein		4.25	
	129506	AB020684	Hs.11217	KIAA0877 protein	6.50		
20	129595	U09550	Hs.1154	oviductal glycoprotein 1, 120kD (mucin 9		10.00	
	130160	AA305688	Hs.267695	UDP-Gal:betaGlcNAc beta 1,3-galactosyltr	20.00		
	130340	D82326	Hs.239106	solute carrier family 3 (cystine, dibasi	11.50		•
	131220	AB023194	Hs.300855	KIAA0977 protein	17.50		
	131430	Al879148	Hs.26770	fatty acid binding protein 7, brain	6.10		
25	132114	NM_006152	Hs.40202	lymphoid-restricted membrane protein		6.15	
	132458	AA935315	Hs.48965	Homo sapiens cDNA: FLJ21693 fis, clone C		5.58	
	132647	NM_006927	Hs.54432	sialyitransferase 4B (beta-galactosidase	7.50		
	132655	D49372	Hs.54460	small inducible cytokine subfamily A (Cy		2.53	
	132682	A1077500	Hs.54900	serologically defined colon cancer antig		2.50	
30	132747	AA345241	Hs.55950	ESTs, Weakly similar to KIAA1330 protein		2.83	
	132812	R50333	Hs.92186	Leman coiled-coil protein		3.82	
	133337	AF085983	Hs.293676	ESTs		5.00	
	133876	AL134906	Hs.771	phosphorylase, glycogen; liver (Hers dis		3.00	
	134119	AW157837	Hs.79226	fasciculation and elongation protein zet		2.06	
35	134464	AA302983	Hs.239720	CCR4-NOT transcription complex, subunit		2.27	•
	134542	M14156	Hs.85112	insulin-like growth factor 1 (somatomedi		11.50	
	135002	AA448542	Hs.251677	G antigen 7B	87.00	0.70	
	135305	AA203555	Hs.98288	Homo sapiens cDNA FLJ14903 fis, clone PL		6.50	
							•

TABLE 6B show the accession numbers for those primekeys lacking unigenelD's for Table 6A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers

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Pkey CAT number Accessions AA100796 AF020589 AA074629 AA075946 AA100849 AA085347 AA126309 AA079311 AA079323 AA085274 X98266 N41124 108562 **55** 103439 35330_1 123551 genbank_AA608837 AA608837 genbank_AA620840 entrez_U92015 AA620840 123861 102832 U92015 101972 entrez_S82472 S82472 60 121558 genbank_AA412497 AA412497

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Table 7A shows 98 genes down-regulated in non-smokers with lung cancer relative to smokers with lung cancer. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probesel obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5

Unique Eos probeset identifier number
Exemplar Accession number, Genbank accession number
Unigene number
Unigene number
Unigene gene title
90th percentile of AI for samples from smokers with adenocarcinoma divided by the average of AI for samples from non-smokers with adenocarcinoma.
90th percentile of AI for samples from smokers with squamous cell carcinoma divided by the average of AI for samples from non-smokers with squamous cell Pkey: ExAccn: UnigenelD: Unigene Title: R1: R2: 10

		carcinor	na.			
	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2
15	100187 100380	D17793 D82343	Hs.78183 Hs.18551	aldo-keto reductase family 1, member C3 neuroblastoma (nerve tissue) protein		164.10 77.40
	100576	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid	102.40	
	100971 101046	BE379727 K01160	Hs.83213	fatty acid binding protein 4, adipocyte (NONE)	463.80 672.00	
20	101066	AW970254	Hs.889	Charot-Leyden crystal protein	66.00	
	101175	U82671	Hs.36980	melanoma antigen, familiy A, 2		77.20
	101497 101663	W05150 NM_003528	Hs.37034 Hs.2178	homeo bax A5	62.80 78.00	
	101663	NM_000715	Hs.1012	H2B histone family, member Q complement component 4-binding protein,	186.20	
25	101745	M88700	Hs.150403	dopa decarboxylase (aromatic L-amino acl	80.08	
	101941	S77583		gb:HERVK10/HUMMTV reverse transcriptase	99.20	400.40
	102125 102242	NM_006456 U27185	Hs.288215 Hs.82547	sialyltransferase retinoic acid receptor responder (tazaro	67.00	103.10
	102340	U37055	Hs.278657	macrophage stimulating 1 (hepatocyte gro	71.60	
30	102369	U39840	Hs.299867	hepatocyte nuclear factor 3, alpha		69.70
	102457	NM_001394	Hs.2359	dual specificity phosphatase 4	153.00	66.70
	102669 102796	U71207 AL079646	Hs.29279 Hs.107019	eyes absent (Crosophila) homolog 2 symplekin; Huntingtin interacting protei		65.70 58.80
	102829	NM_006183	Hs.80962	neurotensin		268.80
35	103207	X72790		gb:Human endogenous retrovirus mRNA for	70.00	
	103242 103260	X76342 X78416	Hs.389 Hs.3155	alcohol dehydrogenase 7 (class IV), mu o casein, alpha		212.10 130.70
	103200	X89211	По. Ј (ОО	gb:H.sapiens DNA for endogenous retrovir	64.60	130.10
40	104212	AB002298	Hs.173035	KIAA0300 protein	66.80	
40	104252	AF002246	Hs.210863	cell adhesion molecule with homology to	63.80	
	104258 105024	AF007216 AA126311	Hs.5462 Hs.9879	solute carrier family 4, sodium bicarbon ESTs	94.40 68.20	
	106260	Al097144	Hs.5250	ESTs, Weakly similar to ALU1_HUMAN ALU S	00.20	74.60
15	106440	AA449563	Hs.151393	glutamate-cystelne ligase, catalytic sub		71.10
45	106566 106605	BE298210	Un 21402	gb:601118016F1 NIH_MGC_17 Homo sapiens c	73.20 83.80	
	106614	AW772298 AA648459	Hs.21103 Hs.335951	Horno sapiens mRNA; cDNA DKFZp5648076 (fr hypothetical protein AF301222	03.00	62.30
	106654	AW075485	Hs.286049	phosphoserine aminotransferase		202.40
50	106999	H93281	Hs.10710	hypothetical protein FLJ20417		89.60
20	108700 108810	AA121518 AW295647	Hs.193540 Hs.71331	ESTs, Moderately similar to 2109260A B c hypothetical protein MGC5350		66.40 95.50
	108857	AK001468	Hs.62180	anillin (Drosophila Scraps homolog), act		63.40
	109597	AA989362	Hs.293780	ESTs	85.00	
55	109691	T65568 A1743680	Hs.12860 Hs.12876	ESTs ESTs		58.70 60.60
55	109704 110942	R63503	Hs.28419	ESTs	76.40	00.00
	111722	R23924	Hs.23596	EST	74.60	
	112891	T03927	Hs.293147	ESTs, Moderately similar to A46010 X-II	64.80	70.70
60	113073	AL157425 N39342	Hs.133315 Hs.103042	Homo sapiens mRNA; cDNA DKFZp761J1324 (f microtubule-associated protein 1B		76.70 120.20
•	114251	H15261	Hs.21948	ESTs	127.20	
	115230	AA278300	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L	174.00	04.00
	115291 115815	BE545072 AW905328	Hs.122579 Hs.180842	hypothetical protein FLJ 10461 ribosomal protein L13	66.40	91.00
65	115909		Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH		226.60
	115965	AA001732	Hs.173233	hypothetical protein FLJ 10970	82.80	004.00
	116107 116552	AL133916 D20508	Hs.172572 Hs.164649	hypothetical protein FLJ20093 hypothetical protein DKFZp434H247	69.00	361.60
	116571	D45652	113.104043	gb:HUMGS02848 Human adult lung 3' direct	64.20	
70	118466	N66741		gb:yz33g08.s1 Morton Fetal Cochlea Homo		63,50
	120484	AA253170	Hs.96473	EST EST	81.60	81.10
	120983 121034	AA398209 AL389951	Hs.97587 Hs.271623	nucleoporin 50kD		66.20
~-	121423	AW973352	Hs.290585	ESTs	64,40	, ,,,,,,,
75	122553	AA451884	Hs.190121	ESTs		60.40
	122946 123130	A1718702 AA487200	Hs.308026	major histocompatibility complex, class gb:ab19f02.s1 Stratagene lung (937210) H	188.60	80.20
	124472	N52517	Hs.102670	EST	71.00	04.E0
90	124526	N62096	Hs.293185	ESTs, Wealdy similar to JC7328 amino aci		104.90
80	125489 125731	H49193	Hs.124984	ESTs, Moderately similar to ALU7_HUMAN A ESTs		72.00 69.90
	· 125747	R61771 NM_002884	Hs.26912 Hs.865	RAP1A, member of RAS oncogene family	69.00	03.30
	126020	H79863	Hs.114243	ESTs		62.40
85	126547	U47732	Hs.84072	transmembrane 4 superfamily member 3		62.80
05	126966	R38438	Hs.182575	solute carrier family 15 (H+/peptide tra		60.10

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		AA761378	Hs.192013	ESTs	70.20		
	127610	AA960867	Hs.150271	ESTs, Highly similar to unnamed protein	64.00		
	127742	AW293496	Hs.180138	ESTs	85.20		
_	127987	AI022103	Hs.124511	ESTs	96.60		
5	128233	AW889132	Hs.11916	ribokinase		78.90	
_	128420	AA650274	Hs.41296	fibronectin leucine rich transmembrane p		106.90	
	128766	AW160432	Hs.296460	craniofacial development protein 1	66.80		
	129014	AW935187	Hs.170162	KIAA1357 protein		58.53	
	129215	AB040930	Hs.126085	KIAA1497 protein	64.20		
10	130090	H97878	Hs.132390	zinc finger protein 36 (KOX 18)	63.80		
	130385	AW067800	Hs.155223	stanniocalcin 2		139.60	
	130732	AW890487	Hs.63984	cadherin 13, H-cadherin (heart)		64.60	
	131025	AB040900	Hs.6189	KIAA1467 protein	64.40		
	131241	BE501914	Hs.24654	Homo sapiens cDNA FLJ11640 fis, clone HE	76.20		
15	131775	AB014548	Hs.31921	KIAA0648 protein	97.80		
1.5	132240	AB018324	Hs.42676	KIAA0781 protein	,	71.00	
	132856	NM_001448	Hs.58367	glypican 4		88.40	
	132977	AA093322	Hs.301404	RNA binding motif protein 3	133.20	00110	
	133749	L20852	Hs.10018	solute carrier family 20 (phosphate tran	100.20	59.30	
20	133818	Al110684	Hs.7645	fibrinogen, B beta polypeptide	341.00	00.00	
20	134264	AF149297	Hs.8087	NAG-5 protein	041.00	64.30	_
		M83772	Hs.80876	flavin containing monooxygenase 3		232.53	•
	134265 134346	X84002	Hs.82037	TATA box binding protein (TBP)-associate	66.00	202.00	
			Hs.8262	lysosomal-associated membrane protein 2	00.00	75.80	
25	134395	AA456539				108.30	• .
23	135047	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	71.40	100.00	
	135056	N75765	Hs.93765	lipoma HMGIC fusion partner ADP-ribosylation factor-like 5	70.40		
	135309	Al564123	Hs.42500	ADP-noosylation ractor-like 5	70.40		
30	oligonucle	ectides were de	signed. Gene	ers for those primekeys lacking unigenelD's for Tal clusters were compiled using sequences derived nt Tools (DoubleTwist, Oakland California). The G	from Genbank ES1	Ts and mRNAs. These se	equences were clustered based on seque
	"Accessio	n" column.				•	•
35	Pkey: CAT num Accession	ber: Gene clust	s probeset idea er number accession numi	·			
40	Pkey	CAT number	r Accessions	.			
10	103207 106566	306354 120358_1	X72790 BE298210 AI885095	Al672315 AW086489 BE298417 AA455921 AA90 Al476470 Al287650 Al885299 Al985381 AW5926)2537 BE327124 F 24 AW340136 Al20	R14963 AA085210 AW274 66556 AA456390 Al31081	4273 Al333584 Al369742 Al039658 15 AA484951
	116571	genbank D4		D45652			
45	118466	genbank_N6		N66741			
	101046	entrez_K011					
	101941	entrez_S775					
	103351	entrez_X892		•			
	123130	genbank_AA		AA487200			
50	123130	Acuncus	1707 200	TOTAL EVV			
50							•

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Table 8A shows 1720 genes either up or down-regulated in lung tumors or chronically diseased lung relative to a broad collection of over 40 distinct normal body tissues. Chronically diseased lung samples represent chronic non-malignant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 39494 probesets on the Eos/Affymetrix Hu02 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5

Pkey: Unique Eos probeset identifier number

Exacch: Exemplar Accession number, Genbank accession number
UnigenelD: Unigene number
Unigene Title: Unigene gene title
R1: 70th percentile of AI for ung tumors divided by 90th percentile of AI for normal lung
R2: 70th percentile of AI for chronically diseased lung divided by 90th percentile of AI for normal lung 10

	RZ:	70th perc	entitle of All for o	stronically diseased lung divided by soru percentile o	ALIOT NOM	iai iung
	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
15	300097	Al916973	Hs.213603	ESTs	5.46	4.69
13	300117	AW189787	Hs.147474	ESTs	0.58	0.56
	300197	AI686661	Hs.218286	ESTs	4.26	5.44
	300201	Al308300		gb:ta90c06.x1 NCI_CGAP_Bm20 Homo sapien	0.62	0.83
	300225	Al989963	Hs.197505	ESTs	1.68	1.75
20	300247	AW274682	Hs.161394	ESTs	1.08	2.28
	300256	A1469095	Hs.298241	Transmembrane protease, serine 3	0.86	1.00
	300337	A1707881	Hs.202090	ESTs	5.80	9.09
	300362	Z42308	11- 04 44 50	gb:HSC0FB121 normalized infant brain cDN	4.18 2.99	12.78 4.38
25	300374	A1859947	Hs.314158	ESTs	1.50	2.53
23	300387 300440	AW270150	Hs.254516 Hs.146164	ESTs ESTs	3.98	5.25
	300440	Al421541 R10367	Hs.307921	EST, Weakly similar to Z232_HUMAN ZINC F	3.18	6.80
	300449	Al362967	Hs.132221	hypothetical protein FLJ12401	0.43	0.62
	300469	AW135830	Hs.233955	hypothetical protein FLJ20401	0.16	0.83
30	300552	X85711	Hs.21838	hypothetical protein FLJ11191	4.10	9.75
	300627	W27363		gb:ab37d01.r1 Stratagene HeLa cell s3 93	4.60	12.60
	300630	AW118822	Hs.128757	ESTs	2.91	5.86
	300716	Al216113	Hs.126280	hypothetical protein FLJ23393	1.00	0.92
25	300738	Al623332	Hs.130541	KIAA1542 protein	1.82	1.71
-35	300777	AA235361	Hs.96840	KIAA1527 protein	4.48	8.22 1.18
	300790	AI492471	Hs.188270	ESTs	1.29 5.51	8.56
	300832	AI688147	Hs.220615 Hs.22958	ESTs, Weakly similar to T03829 transcrip calcium channel alpha2-delta3 subunit	4.90	6.34
	300836 300838	Z44942 Al582897	Hs.192570	hypothetical protein FLJ22028	1.70	2.81
40	300878	AW449802	Hs.285901	Homo sapiens cDNA FLJ20428 fis, clone KA	4.56	7.91
-10	300897	Al890356	Hs.127804	ESTs, Weakly similar to T17233 hypotheti	2.23	1.58
	300926	AA504860		gb:ab03a10.s1 Stratagene fetal retina 93	2.13	3.50
	300960	AI041019	Hs.152454	ESTs	2.74	4.46
	300961	AW204069	Hs.312716	ESTs, Weakly similar to unnamed protein	1.00	1.00
45	300962	AA593373	Hs.293744	ESTs	1.46	1,51
	300967	AA565209	Hs.269439	ESTS	0.39 1.49	1.30 1.08
	300987	AW450840	Hs.148590	ESTs, Weakly similar to AF208846 1 BM-00	0.16	0.37
	300988 301050	Al927208 AW136973	Hs.208952 Hs.288516	ESTs ESTs, Weakly similar to S69890 mitogen i	3.23	1.94
50	301098	AA677570	Hs.185918	ESTs	6.76	14.28
50	301157	AA729905	Hs.231916	ESTs	3.16	8.85
	301162	AI142118	Hs.129004	ESTs	1.68	7.18
	301170	AA737594	Hs.247606	ESTs	4.40	6.42
	301192	AI808751	Hs.121188	ESTs	6.38	11.59
55	301193	AA758115	Hs.128350	ESTs, Weakly similar to JC5423 2-hydroxy	4.35	7.78
	301267	AW297762	Hs.255690	ESTs	1.56	1.61
	301281	AA843986	Hs.190586	ESTs	2.19 0.76	1.78 0.76
	301341	AI819198	Hs.208229	ESTs .	1.00	1.81
60	301382 301407	AA912839 AW450466	Hs.163369 Hs.126830	ESTs	1.48	1.51
00	301452	AA975688	Hs.159955	ESTs	0.51	1.46
	301483	AW272467	Hs.254655	Untitled	2.40	5.02
	301494	A1678034	Hs.131099	ESTs	2.79	3.41
	301521	A1733621	Hs.133011	zinc finger protein 117 (HPF9)	0.67	0.67
65	301531	Al077462	Hs.134084	ESTs	2.52	3.76
	301580	Al878959	Hs.73737	splicing factor, arginine/serine-rich 1	7.41	11.92
	301676	Z43570	Hs.27453	ESTs, Moderately similar to G01251 Rar p	8.31	10.70
	301690	F05865	Hs.108323	ubiquitin-conjugating enzyme E2E 2 (homo DKFZP434F162 protein	2.70 4.20	4.22 8.78
70	301718 301799	F07744 AA384252	Hs.7987 Hs.286132	D15F37 (pseudogene)	5.93	7.04
, 0	301804	AA581004	Hs.62180	antillin (Drosophila Scraps homolog), act	1.70	0.76
	301822	X17033	Hs.271986	Integrin, alpha 2 (CD49B, alpha 2 subuni	1.58	1.36
	301846	R20002	Hs.6823	hypothetical protein FLJ10430	1.00	1.00
	301868	T71508	Hs.13861	ESTs, Weakly similar to pH sensitive max	2.88	5.49
75	301882	T78054		gb:yc97g09.r1 Soares infant brain 1NIB H	2.28	3.80
	301905	A1991127	Hs.117202	ESTs	1.00	1.00
	301948	AA344647	Hs.116724	aldo-kelo reductase family 1, member B11	5.28	2.28
	301960	AW070252	Hs.27973	KIAA0874 protein	5.38	6.48 3.42
80	302011	T91418	Hs.125156	transcriptional adaptor 2 (ADA2, yeast,	3.03 1.00	1.25
90	302016 302041	N40834 NM_001501	Hs.23495 Hs.129715	hypothetical protein FLJ11252 gonadotropin-releasing hormone 2	0.71	0.99
	302072	AJ238381	Hs.132576	paired box gene 9	1.60	1.71
	302094	Al286176	Hs.6786	ESTs	0.52	1.20
0.5	302095	AW044300	Hs.137506	Homo sapiens BAC clone RP11-120J2 from 7	2.75	4.93
85	302148	AW269618	Hs.23244	ESTs	3.04	3.87

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	302155	AI088485	Hs.144759	ESTs	0.45	1.15
	302201	AJ006276	Hs.159003	transient receptor potential channel 6	0.33	0.84
	302202	AF097159	Hs.159140	UDP-Gal:betaGlcNAc beta 1,4- galactosylt	0.52	0.94
5	302206	AI937193	Hs.41143	phosphoinositide-specific phospholipase	2.76	3.65
3	302209 302235	AF047445	Hs.159297 Hs.166361	killer cell lectin-like receptor subfami Homo sapiens mRNA; cDNA DKFZp564F112 (fr	1.00 1.68	1.00 1.50
	302233	AL049987 AL117607	Hs.175563	Homo sapiens mRNA; cDNA DKFZp564N0763 (f	1.00	2.11
	302328	AA354849	Hs.23240	Homo sapiens cDNA FLJ13496 fis, clone PL	9.38	13.08
	302346	AL039101	Hs.194625	dynein, cytoplasmic, light intermediate	3.27	7.24
10	302360	AJ010901	Hs.198267	mucin 4, tracheobronchial	2.54	1.88
	302384	Y08982	Hs.202676	synaptonemal complex protein 2	1.00	0.91
	302406	U86751 AF155156	Hs.211956	CD3-epsilon-associated protein; antisens adaptor-related protein complex 4, epsil	2.63 5.82	2.67 9.34
	302409 302423	AB028977	Hs.218028 Hs.225974	KIAA1054 protein	3.66	3.18
15	302432	AL080068	Hs.272534	Homo sapiens mRNA; cDNA DKFZp564J062 (fr	2.44	6.77
	302435	AF092047	Hs.227277	sine oculis homeobox (Drosophila) homolo	0.44	0.84
	302437	AB024730	Hs.227473	UDP-N-acetylglucosamine:a-1,3-D-mannosid	4.18	5.64
	302455	AA356923	Hs.240770	nuclear cap binding protein subunit 2, 2	1.85	0.92
20	302472	AA317451	Hs.6335 ·	SWI/SNF related, matrix associated, acti U6 snRNA-associated Sm-like protein LSm8	2.04 1.44	2.13 1.89
20	302476 302489	AF182294 T80660	Hs.241578 Hs.230424	Homo saplens cDNA FLJ13540 fis, clone PL	0.51	1.10
	302490	AA885502	Hs.187032	ESTs	2.64	4.87
	302562	AJ005585	Hs.48956	gap junction protein, beta 6 (connexin 3	5.34	2.68
25	302566	AA085996	Hs.248572	hypothetical protein FLJ22965	1.00	1.21
25	302630	AB029488	Hs.272100	SMS3 protein	0.52	1.24
	302634	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophila, mous	1.00 1.58	1.00 1.02
	302638 302647	AA463798 X57723	Hs.102696 Hs.198273	MCT-1 protein NADH dehydrogenase (ubiquinone) 1 bela s	2.72	6.85
	302655	AJ227892	Hs.146274	ESTs	1.00	4.32
30	302656	AW293005	Hs.70704	Homo sapiens, clone IMAGE:2823731, mRNA,	2.97	0.93
	302668	AA580691	Hs.180789	S164 protein	0.80	0.95
	302679	H65022	11 25040	gb:yu66g11.r1 Weizmann Olfactory Epithel	1.68	5.04
	302680	AW192334 AJ001408	Hs.38218	ESTs	2,70 4.25	7.98 8.13
35	302697 302705	U09060		gb:Homo saplens mRNA for immunoglobulin gb:Human immunoglobulin heavy chain, V-r	3.91	8.68
55	302711	L08442		gb:Human autonomously replicating sequen	2.20	2.73
	302719	W69724	Hs.288959	hypothetical protein FLJ20920	0.54	1.02
	302742	L12069		gb:Homo saplens (clone WR4.10VH) anti-th	4.28	11.57
40	302755	AW384815	Hs.149208	KIAA1555 protein	1.57	2.38
40	302771 302789	H98476 AJ245067	Hs.42522	ESTs gb:Homo sapiens mRNA for immunoglobulin	2.94 3.49	4.68 6.31
	302795	AJ245313	Hs.272838	hypothetical protein FLJ 10494	0.80	2.74
	302802	Y08250		gb:H.sapiens mRNA for variable region of	1.13	0.77
	302803	AA442824	Hs.293961	ESTs, Moderately similar to putative DNA	3,14	10.68
45	302812	N31301	Hs.152664	hypothetical protein FLJ20051	3.04	8.24
	302847	X98940	11- 400403	gb:H.sapiens rearranged lg heavy chain (1.80	1.92 1.00
	302885 302943	AL137763 AI581344	Hs.132127 Hs.127812	hypothetical protein LOC57822 ESTs, Wealdy similar to T17330 hypotheti	1.00 0.53	0.67
	302977	AW263124	Hs.315111	hypothetical protein FLJ 12894	2.45	2.62
50	303006	AF078950	Hs.24139	Homo sapiens cDNA: FLJ23137 fis, clone L	4.88	8.61
	303011	AF090405		gb:Homo sapiens clone 2A1 scFV enitbody	1.41	1.86
	303013	F07898	Hs.288968	RAB22A, member RAS oncogene family	1.51	1.19
	303061	AF151882	Hs.27693	peptidylprolyl Isomerase (cyclophilin)-l gb:H.saptens T-cell receptor mRNA	0.72 1.17	0.76 3.90
55	303077 303090	AF163305 AA443259	Hs.146286	kinesin family member 13A	4.08	6.46
<i></i>	303091	AF192913	Hs.130683	zinc finger protein 180 (HHZ168)	2.50	4.37
	303094	AF195513	Hs.278953	Pur-gamma	5.38	8.38
	303095	AF202051	Hs.134079	NM23-H8	3.26	4.08
40	303131	AW081061	Hs.103180	DC2 protein	2.02	1.83
60	303195 303196	AA082211 AA082298	Hs.233936 Hs.59710	myosin, light polypeptide, regulatory, n ESTs	1.32 0.77	3.95 0.53
	303216	AA581439	Hs.152328	ESTs	0.24	0.63
	303222	AA333538	Hs.204501	hypothetical protein FLJ 10534	3.56	6.22
	303234	AA132255	Hs.143951	ESTs	2.28	3.17
65 ·	303251	AW340037	Hs.115897	protocadherin 12	0.38	1.02
	303295	AA205625	Hs.208067	ESTS	2.30	1.00 4.48
	303297 303316	T80072 AF033122	Hs.13423 Hs.14125	Homo sapiens clone 24468 mRNA sequence p53 regulated PA26 nuclear protein	1.86 0.10	0.80
	303467	AA398801	Hs.323397	ESTs	4.54	9.65
70	303506	AA340605	Hs.105887	ESTs, Weakly similar to Homolog of rat Z	0.09	0.04
	303552	AA359799	Hs.224662	ESTs, Weakly similar to unnamed protein	1.00	1.72
	303598	AA382814	11. 04070	gb:EST96097 Testis I Homo sapiens cDNA 5	4.96	9.14
	303637	AF056083	Hs.24879	phosphatidic acid phosphatase type 2C	2.06 1.00	2.02 1.24
75	303655 303756	AA504702 A1738488	Hs.258802 Hs.115838	ATPase, (Na+)/K+ transporting, beta 4 po ESTs	1.08	1.43
. •	303856	AA968589	Hs.180532	glucose phosphate isomerase	1.76	1.31
	303893	N88597	Hs.113503	karyopherin (importin) beta 3	2.30	2.57
	303907	AW467774	Hs.171880	polymerase (RNA) II (DNA directed) polyp	3.10	5.79
QΛ	303946	AW474196	Hs.306637	Homo sapiens cDNA FLJ12363 fis, clone MA	5.06	11.86
80	303978 303981	AW513315 AW513804	Hs.278834	gb:xo43c12.x1 NCI_CGAP_Ut1 Homo sapiens ESTs, Weakly similar to ALU1_HUMAN ALU S	5.14 2.83	7.31 4.06
	303990	AW515465	113.21 0034	gb:xu71a11.x1 NCI_CGAP_Kld8 Homo sapiens	1.15	2.35
	303998	AW516449		gb:xl68f05.x1 NCI_CGAP_Ut2 Homo sapiens	2.20	9.35
0.5	303999	AW516611		gb:xp70b11.x1 NCI_CGAP_Ov39 Homo sapiens	4.85	6.28
85	304006	AW517947		gb:xt66h02.x1 NCI_CGAP_Ut2 Homo sapiens	3.21	4.07

	w	O 02/08	6443			
	304008	AW518198	Hs.3297	ribosomal protein S27a	6.50	11.08
	304009	AW518206	Hs.181165	eukaryolic translation elongation factor	1.88	3.27
	304024	T03036		gb:FB21B7 Fetal brain, Stratagene Homo s	2.15	3.55
_	304026			gb:FB26F2 Fetal brain, Stratagene Homo s	5.88	11.80
5	304028	T03266		gb:FB7C1 Fetal brain, Stratagene Homo sa	5.59	13.46
	304036	T16855	Hs.244621	ribosomal protein S14	6.55	14.43
	304046	T54803		gb:yb42d06.s1 Stratagene fetal spleen (9	6.18 2.64	12.19 8.23
	304061 304063	T61521 T62536		gb:yb73g01.s1 Stratagene ovary (937217) gb:yc04c12.s1 Stratagene lung (937210) H	0.53	1.61
10	304097	R25376	Hs.177592	ribosomal protein, large, P1	6.49	11.67
	304114	R78946		gb:yl87g02.s1 Soares placenta Nb2HP Homo	2.90	4.18
	304122	H28966		gb:ym31a06.s1 Soares infant brain 1NIB H	1.00	2.76
	304155	H68696		gb:yr78b06.s1 Soares fetal liver spleen	0.79	1.18
1.5	304203	N56929		gb:yy82d08.s1 Soares_multiple_sclerosis_	4.28	11.34
15	304234	W81608	11- 20240	gb:zd88h06.s1 Soares_fetat_heart_NbHH19W	6.47	11.03 1.16
	304267	AA064862	Hs.73742 Hs.297753	ribosomal protein, large, P0 vimentin	1.34 3.40	5.40
	304270 304287	AA069711 AA079286	Hs.78466	proteasome (prosome, macropain) 26S sub	2.93	4.42
	304348	AA179868	113.7 0400		3.98	10.96
20	304415	AA290747	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	3.32	5.99
	304430	AA347682		gb:EST54044 Fetal heart II Homo sapiens	1.00	1.00
	304456	AA411240		gb:zv26g05.s1 Soares_NhHMPu_S1 Homo sapi	1.42	3.33
	304521	AA464716		gb:zx82c11.s1 Soares ovary tumor NbHOT H	2.18	1.15
25	304526	AA476427		gb:zx02c05.s1 Soares_total_fetus_Nb2HF8_	5.38	14,11
25	304542	AA482602	Hs.169476	glycereldehyde-3-phosphate dehydrogenase	4.16	8.23 1.20
	304546	AA486074	Hs.297681	serine (or cysteine) proteinase inhibito gb:nh85e08.s1 NCI_CGAP_Br1.1 Homo saplen	0.55 1.95	2.10
	304607 304640	AA513322 AA524440	Hs.111334	ferritin, light polypeptide	2.10	2.83
	304650	AA527489	Hs.3463	ribosomal protein S23	3.33	12.62
30	304735	AA576453	113.0400	gb:nm75h11.s1 NCI_CGAP_Co9 Homo sapiens	1.33	0.88
	304760	AA580401		gb:nn13g09.s1 NCI_CGAP_Co12 Homo saplens	3.68	8.14
	304849	AA588157	Hs.13801	KIAA1685 protein	2.77	3.70
	304917	AA602685	Hs.284136	PRO2047 protein	7.16	11.01
25	304921	AA603092	Hs.297753	vimentin	2.47	4.24
35	304966	AA613893	Hs.282435	ESTs .	6.78	11.66
	304987	AA618044	Hs.300697	Immunoglobulin heavy constant gamma 3 (G	0.90 6.46	1.23 10.17
	305016	AA626876		gb:zu89h06.s1 Soares_testis_NHT Homo sap gb:ab99c04.s1 Stratagene lung (937210) H	1.00	1.00
	305034 305072	AA630128 AA641012		gb:nr72a12.s1 NCI_CGAP_Pr24 Homo sapiens	5.68	11.59
40	305111	AA644187	Hs.303405	ESTs	1.48	1.37
10	305148	AA654070	113.000 700	gb:nt01g08.s1 NCI_CGAP_Lym3 Homo sapiens	1.76	4.61
	305159	AA659166	Hs.275668	EST, Weakly similar to EF1D_HUMAN ELONG	1.00	2.15
	305190	AA665955		gb:ag57d12.s1 Gessler Wilms tumor Homo s	5.31	8.14
4.5		AA670052	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	0.78	1.18
45	305235	AA670480		gb:ag37e01.s1 Jia bone marrow stroma Hom	3.11	8.66
	305245	AA676695	Hs.81328	nuclear factor of kappa light polypeptid	4.38	7,53
	305312 305322	AA700201	Un 162010	gb:zj44f07.s1 Soares_fetal_liver_spleen_ EST	2.13 1.20	2.66 1.40
	305322	AA701597 AA720942	Hs.163019 Hs.300697	immunoglobulin heavy constant gamma 3 (G	1.16	0.68
50	305413	AA724659	113.300031	gb:ai10f08.s1 Soares_parathyroid_tumor_N	5.86	9.87
	305447	AA737856		gb:nx10c08.s1 NCI_CGAP_GC3 Homo sapiens	2.21	2.86
	305476	AA745664	Hs.287445	hypothetical protein FLJ11726	3.36	6.54
	305483	AA748030	Hs.303512	EST	1.00	2.02
	305528	AA769156		gb:nz12e05.s1 NCI_CGAP_GCB1 Homo sapiens	6.44	9.10
55	305612	AA782347	Hs.272572	hemoglobin, alpha 2	0.19	0.79
	305614	AA782866	11- 03F00F	gb:aj09h02.s1 Soares_parathyroid_tumor_N	1.00	1.00
	305616	AA782884	Hs.275865	ribosomal protein S18 gb:ce29a12.s1 NCI_CGAP_Pr25 Homo sapiens	7.57 4.78	10.20 12.42
	305637 305639	AA806124 AA806138		gb:oe29c12.s1 NCI_CGAP_Pr25 Homo sapiens	0.89	0.70
60	305650	AA807709		gb:nw31e04.s1 NCI_CGAP_GCB0 Homo sapiens4		8.71
	305690	AA813477		gb:ai67a05.s1 Soares_testis_NHT Homo sap	4.91	9.40
	305726	AA828156	Hs.73742	ribosomal protein, large, P0	0.19	0.81
	305728	AA828209		gb:of34a02.s1 NCI_CGAP_Kid6 Homo sapiens	5.12	9.29
65	305759	AA835353		gb:ak72b06.s1 Barstead spleen HPLRB2 Hom	1.66	4.11
65	305792	AA845256	11. 70740	gb:ak84a08.s1 Barstead spleen HPLRB2 Hom	2.34	4.25
	305864	AA864374	Hs.73742	ribosomal protein, large, P0	0.30 2.10	1.40 5.21
	305901 305910	AA872968 AA875981		gb:oh63h08.s1 NCI_CGAP_Kid5 Homo sapiens gb:nx21h02.s1 NCI_CGAP_GC3 Homo sapiens	0.32	1.01
	306015	AA897116		gb:am08b07.s1 Soares_NFL_T_GBC_S1 Homo s1		1.12
70	306017	AA897221	Hs.109058	ribosomal protein S6 kinase, 90kD, polyp	5.21	7.90
	306020	AA897630	Hs.130027	EST	1.96	6.59
	306063	AA906316		gb:ok03g03.s1 Soares_NFL_T_GBC_S1 Homo s	7.38	20.69
	306065	AA906725		gb:ok78g02.s1 NCI_CGAP_GC4 Homo saplens	7.19	13.48
75	306104	AA910956		gb:ok85h11.s1 NCI_CGAP_Kid3 Homo sapiens	6.50	9.13
75	306109	AA911861	11- 000000	gb:og21a07.s1 NCI_CGAP_PNS1 Homo sapiens	4.21	5.25
	306148	AA917409	Hs.288036	tRNA isopentenylpyrophosphate transferas	2.20 2.84	2.70 5.35
	306242	AA932805		gb:oo60g04.s1 NCI_CGAP_Lu5 Homo sapiens gb:oi53h05.s1 NCI_CGAP_HN3 Homo sapiens	1.60	1.12
	306288 306325	AA936900 AA953072	Hs.210546	interleukin 21 receptor	1.65	2.26
80	306353	AA961382	Hs.275865	ribosomal protein \$18	3.78	6.32
	306375	AA968650	Hs.276018	EST, Moderately similar to JC4662 ribos	4.30	5.74
	306396	AA970223		gb:op09d05.s1 NCI_CGAP_Kid6 Homo saplens	0.95	2.45
	306428	AA975110	Hs.191228	hypothetical protein FLJ20284	3.19	4.10
0.5	306442	AA976899		gb:oq35e09.s1 NCI_CGAP_GC4 Homo sapiens	4.67	7.44
85	306446	AA977348		gb:oq72e12.s1 NCI_CGAP_Kid6 Homo sapiens	3.92	6.27

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	306458	AA978186		gb:op33c06.s1 Soares_NFL_T_GBC_S1 Homo s	3.35	5.77
	306467	AA983508	Hs.163593	ribosomal protein L18a	3.72	5.37
	306510	AA988546		gb:or84d07.s1 NCL_CGAP_Lu5 Homo saplens	1.00	1.00
5	306555	AA994304	Hs.276083	EST, Weakly similar to RL23_HUMAN 60S R gb:ou57e08.s1 NCI_CGAP_Br2 Homo sapiens	6.61 16.20	10.91 31.83
5	306557 306572	AA994530 AA995686		gb:0057e00.81 NCI_CGAP_612 Horno sapiens gb:0525c12.91 NCI_CGAP_Kid5 Horno sapiens	2.51	6.52
	306582	AA996248		gb:os18c10.s1 NCI_CGAP_Kld5 Homo sapiens	1.42	3.13
	306598	A1000320	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	4.91	8.68
10	306605	A1000497	Hs.119500	ribosomal protein, large P2	1.96	8.60
10	306656	AI004024	11- 204420	gb:ou11b07.x1 Soares_NFL_T_GBC_S1 Homo s	0.11 9.56	0.45 17.28
	306676 306686	A1005603 A1015615	Hs.284136	PRO2047 protein gb:ov29f10.x1 Soares_testis_NHT Homo sap	1.86	3.60
	306702	AI022565	Hs.307670	EST COMES_COSS_CONT FISHER COSS	1.47	1.19
	306728	AI027359	Hs.272572	hemoglobin, alpha 2	1.28	2.83
15	306751	AI032589		gb:ow70h12.s1 Soares_fetal_liver_spleen_	3.91	5.21
	306767	AI038963	Hs.249118	ESTS	3.33 3.77	6.06 7.46
	306892 306897	Al092465 Al093967		gb:qa75h12.x1 Soares_fetal_heart_NbHH19W gb:qa33c06.s1 Soares_NhHMPu_S1 Homo sapi	2.12	2.85
	306956	A)125111		gb:am66f03.s1 Barstead spleen HPLRB2 Hom	6.10	10.52
20	306958	A)125152		gb:am55e09.x1 Johnston frontal cortex Ho	1.72	1.56
	307035	Al142774	Hs.119122	ribosomal protein L13a	2.00	4.70
	307041	AI144243		gb:qb85b12.x1 Soares_fetal_heart_NbHH19W	9.12	12.56 8.52
	307091	AI167439		gb:ox70h06.s1 Soares_NhHMPu_S1 Homo sapi gb:qc99g06.x1 Soares_pregnant_uterus_NbH	4.88 3.55	6.44
25	307181 307297	Al189251 Al205798	Hs.111334	ferritin, light polypeptide	2.46	4.65
	307317	A1208303	Hs.147333	EST	5.64	10.13
	307327	Al214142	Hs.246381	CD68 antigen	3.18	5.15
	307382	Al223158	Hs.147885	ESTs	2.02	3.73
20	307410	AI241715	Hs.77039	ribosomal protein S3A gb:qh92b02.x1 Soares_NFL_T_GBC_S1 Homo s	0.72 2.38	0.48 3.51
30	307415 307423	Al242118 Al243206	Hs.179573	collagen, type I, alpha 2	2.60	5.44
	307426	A1243364	115.11.0010	gb:gh30g11.x1 Soares_NFL_T_GBC_S1 Homo s	3.18	7.67
	307517	A1275055		gb:gl72d03.x1 Soares_NhHMPu_S1 Homo sapi	1.00	1.00
25	307551	AI281556		gb:qu52f11.x1 NCI_CGAP_Lym6 Homo sapiens	3.40	11.20
35	307561	A1282207		gb:qp65a12.x1 Soares_fetal_lung_NbHL19W	4.74 3.50	15.51 7.19
	307608 307657	Al290295 Al306428	Hs.298262	gb:qm01f02.x1 Soares_NhHMPu_S1 Homo sapi ribosomal protein S19	1.76	2.44
	307691	Al318285	113.230202	gb:tb17b01.x1 NCI_CGAP_Ov37 Homo sapiens	1.59	1.31
	307701	A1318583	Hs.276672	EST, Weakly similar to RL6_HUMAN 60S RI	1.90	2.13
40	307718	Al333406	Hs.83753	small nuclear ribonucleoprotein polypept	0.45	0.99
	307730	Al336092		gb:qt43b07.x1 Soares_fetal_lung_NbHL19W	1.51 1.00	0.99 1.00
	307760 307764	Al342387 Al342731		gb:qt27f07.x1 Soares_pregnant_uterus_NbH gb:qo26a07.x1 NCI_CGAP_Lu5 Homo sapiens	4.52	12.58
	307783	A1347274		gb:tc05d02.x1 NCI_CGAP_Co16 Homo sapiens	1.42	1.00
45	307796	Al350556		gb:qt18f09.x1 NCI_CGAP_GC4 Homo sapiens	6.57	9.61
	307807	Al351799		gb:qt09d02.x1 NCI_CGAP_GC4 Homo sapiens	3.38	7.68 0.86
	307808	AI351826		gb:qt09g03.x1 NCI_CGAP_GC4 Homo sapiens gb:qt94a11.x1 NCI_CGAP_Co14 Homo sapiens	0.33 7.94	21.57
	307820 307830	Al355761 Al358722	Hs.276737	EST, Weakly similar to RSHU22 ribosomal	2.05	3.32
50	307852	Al365541	110.12.1 01 07	gb:qz08g05.x1 NCI_CGAP_CLL1 Homo sapiens	3.18	5.21
	307902	Al380462		gb:tg02h05.x1 NCI_CGAP_CLL1 Homo sapiens	3.13	4.99
	307997	AI434512	Hs.181165	eukaryotic translation elongation factor	1.00	3.01 12.64
	308002 308011	AI435240	Hs.283442	ESTs gb:ti60a08.x1 NCI_CGAP_Lym12 Homo sapien	5.86 3.79	5.83
55	308023	A1439473 A1452732	Hs.251577	hemoglobin, alpha 1	0.38	0.88
-	308041	AI458824	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	4.36	6.06
	308059	AI468938	Hs.276877	EST, Weakly similar to RL10_HUMAN 60S R	1.80	1.98
	308085	AJ474135	Hs.181165	eukaryotic translation elongation factor	3.38 1.30	4.14 3.87
60	308101	A1475950	Hs.181165	eukaryotic translation elongation factor gb:tj77e12.x1 Soares_NSF_F8_9W_OT_PA_P_S2		8.72
00	308106 308122	A1476803 A1480123	Hs.309411	EST	2.70	3.86
	308154	A1500600		gb:tn93d08.x1 NCI_CGAP_Ut2 Homo sapiens	0.66	1.33
	308171	A1523632	Hs.298766	ESTs, Weakly similar to schlafen4 [M.mu	2.48	4.86
65	308211	Al557029	Hs.278572	anaplastic lymphoma kinase (Ki-1)	2.43 3.34	2.14 3.79
65	308213 308216	Al557041 Al557135		gb:PT2.1_12_E04.r tumor2 Homo sapiens cD gb:PT2.1_13_H06.r tumor2 Homo sapiens cD	4.61	4.78
	308219	AI557246		gb:PT2.1_15_D07.r tumor2 Homo sapiens cD	4.87	7.94
	308271	AI567844	Hs.252259	ribosomal protein S3	2.40	6.35
70	308319	AI583983	Hs.181165	eukaryotic translation elongation factor	2.45 .	3.33
70	308362	A1613519	Hs.105749	KIAA0553 protein	1.24 3.16	1.41 4.82
	308413 308450	Al636253 Al660860	Hs.196511 Hs.96840	ESTs KIAA1527 protein	1.79	2.68
	308464	A1672425	Hs.277117	EST, Moderately similar to 138055 myosi	4.87	8.27
	308588	AJ718299		gb:as51g12.x1 Barstead aorta HPLRB6 Homo	3.90	5.64
75	308599	AI719893		gb:as47d07.x1 Barstead aorta HPLRB6 Homo	3.32	5.12
	308615	A1738593	Hs.101774	hypothetical protein FLJ23045	3.11 3.98	2.36 3.69
	308643 308673	A1745040 A1760864		gb:tr19a12.x1 NCI_CGAP_Ov23 Homo sapiens gb:wi09c10.x1 NCI_CGAP_CLL1 Homo sapiens	0.82	0.99
<u>.</u> .	308697	Al760004 Al767143		gb:wi97a07.x1 NCI_CGAP_KId12 Homo sapien	2.76	5.59
80	308762	AI807405	Hs.259408	ESTs	3.17	6.30
	308778	AI811109	11- 646-	gb:tr04c11.x1 NCI_CGAP_Ov23 Homo saplens	1.00	1.00
	308782	AJ811767	Hs.2186	eukaryotic translation elongation factor gb:wk52c01.x1 NCl_CGAP_Pr22 Homo saptens	2.94 4.41	5.15 8.34
	308808 308823	AI818289 AI824118	Hs.217493	annexin A2	1.85	1.92
85	308875	AI832332		gb:at48g03.x1 Barstead colon HPLRB7 Homo	2.52	3.80

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	308879	AI832763	Hs.75968	thymosin, beta 4, X chromosome	3.38 3.06	7.96 2.65
	308886 308898	A1833240 A1858845		gb:at76d10.x1 Barstead colon HPLRB7 Homo gb:wt32d10.x1 NCI_CGAP_Ut1 Homo saptens	2.45	3.44
	308934	A1865023	Hs.177	phosphatidylinositol glycan, class H	4.14	6.76
5	308966	AI870704		gb:wl47h01.x1 NCI_CGAP_Ut1 Homo saplens	1.00	1.00
	308979	Al873111		gb:w/52h05.x1 NCI_CGAP_Bm25 Homo sapien	7.15 0.61	11.10 0.59
	309045 309051	Al910902 Al911975		gb:tq39f01.x1 NCI_CGAP_Ut1 Home sapiens gb:wd78d01.x1 NCI_CGAP_Lu24 Home sapiens	1.78	4.42
	309069	Al917366	Hs.78202	SWI/SNF related, matrix associated, act	3.27	5.88
10	309083	Al922426	Hs.119598	ribosomal protein L3	2.39	3.34
	309105	A1925503	Hs.265884	ESTS	5.54 1.00	17.78 2.92
	309122 309128	Al928178 Al928816	Hs.180842	gb:wo95a11.x1 NCI_CGAP_Kid11 Homo saplen ribosomal protein L13	1.38	5.55
	309164	AI937761	113.100072	gb:wp84b09.x1 NCI_CGAP_Bm25 Homo sapien	2.43	3.11
15	309177	Al951118		gb:wx63g05.x1 NCI_CGAP_Br18 Homo sapiens	0.81	0.97
	309288	Al991525	Hs.299426	ESTS	4.86 4.36	7.46 9.43
	309299 309303	AW003478 AW004823		gb:wq66c06.x1 NCI_CGAP_GC6 Homo sapiens gb:ws93a08.x1 NCI_CGAP_Co3 Homo sapiens	2.88	7.54
	309303	AW085201	Hs.244144	EST	4.30	7.14
20	309437	AW090702	Hs.278242	tubulin, alpha, ubiquitous	2.49	3.11
	309459	AW117645	Hs.65114	keratin 18	2.88	4.55
	309476 309499	AW129368 AW136325	Hs.279771	gb:xe14b05.x1 NCI_CGAP_Ut4 Homo sapiens Homo sapiens clone PP1596 unknown mRNA	2.08 2.82	6.60 3.55
	309529	AW150807	Hs.181357	taminin receptor 1 (67kD, ribosomal pro	4.78	3.95
25	309532	AW151119		gb:xg33e10.x1 NCt_CGAP_Ut1 Homo sapiens	1.18	4.40
	309626	AW192004	Hs.297681	serine (or cysteine) proteinase inhibit	4.46 1.47	12.06 1.39
	309641 309675	AW194230 AW205681	Hs.253100 Hs.253506	EST, Moderately similar to GHHU Ig gamm EST, Moderately similar to ATPN_HUMAN A	5.68	15.20
	309693	AW237221	Hs.181357	laminin receptor 1 (67kD, ribosomal prot	1.00	1.00
30	309695	AW238011	Hs.295605	mannosidase, alpha, class 2A, member 2	5.45	9.61
	309700	AW241170	Hs.179661	tubulin, beta polypeptide	1.41	1.25 8.35
	309747 309769	AW264889 AW272346		gb:xq36h02.x1 NCI_CGAP_Lu28 Homo sapiens gb:xs13c10.x1 NCI_CGAP_Kid11 Homo sapien	5.00 5.76	11.90
	309782	AW275156	Hs.156110	immunoglobulin kappa constant	0.42	0.69
35	309783	AW275401	Hs.254798	EST	1.00	4.11
	309799	AW276964		gb:xp58h01.x1 NCI_CGAP_Ov39 Homo sapiens	1.68 3.02	1.44 5.04
	309866 309903	AW299916 AW339071	Hs.300697	gb:xs44c01.x1 NCl_CGAP_Kld11 Homo sapien immunoglobulin heavy constant gamma 3 (G	1.05	1,18
	309923	AW340684	112,000071	gb:hd05g08.x1 Soares_NFL_T_GBC_S1 Homo s	2.30	3.67
40	309928	AW341418		gb:hd08c03.x1 Soares_NFL_T_GBC_S1 Homo s	7.41	13.71
	309931	AW341683		gb:hd13d01.x1 Soares_NFL_T_GBC_S1 Home s	1.20 4.90	12.70 18.29
	309933 309964	AW341936 AW449111	Hs.257111	gb:hb73f10.x1 NCI_CGAP_Ut2 Homo sapiens hypothetical protein MGC3265	1.99	3.07
	310002	AI439096	Hs.323079	Homo sapiens mRNA; cDNA DKFZp564P116 (fr	0.20	0.47
45	310096	AW136822	Hs.172824	ESTs, Weakly similar to B48013 proline-r	1.51	1.22
	310098	AI685841	Hs.161354	ESTs	0.31 2.06	0.76 5.83
	310109 310112	Al203094 AW197233	Hs.148633 Hs.147253	ESTs ESTs	2.92	3.55
	310115	Al611317	Hs.223796	ESTs	1.25	0.84
50	310121	AW195642	Hs.148901	ESTs	1.00	2.71
	310146	A)206614	Hs.197422	ESTS	9.50 2.85	15.31 4.18
	310193 310255	AI627653 AW450439	Hs.147562 Hs.153378	ESTs ESTs	4.26	10.63
	310261	A1240483	Hs.201217	ESTs	3.28	4.40
55	310264	AI915771	Hs.74170	metallothionein 1E (functional)	0.26	0.86
	310275	AJ242102	Hs.213636	ESTs ESTs	5.43 3.15	8.19 8.06
	310282 310290	AJ243332 AW013815	Hs.156055 Hs.149103	ESTs	2.19	3.12
	310333	AJ253200	Hs.145402	ESTs	1,17	1.91
60	310346	AJ261340	Hs.145517	ESTs	4.81	9.95 7.79
	310385 310443	AI263392 AW119018	Hs.156151 Hs.164231	ESTs ESTs	5.96 2.90	4.63
	310444	AW196632	Hs.252956	ESTs	0.85	1.01
<i>-</i> -	310446	AJ275715	Hs.145926	ESTs	2.18	3.85
65	310468	AI984074	Hs.196398	ESTs	3.39 1.00	5.19 1.00
•	310477 310512	AI948801 AW275603	Hs.171073 Hs.200712	ESTs ESTs		8.12
	310512	AI681145	Hs.160724	ESTs	3.30	7.33
70	310524	AW082270	Hs.12496	ESTs, Highly similar to AC004836 1 simil	0.72	1.44
70	310547	AI302654	Hs.208024	ESTs	. 3.26 2.39	3.46 4.08
•	310584 310608	A1653007 A1962234	Hs.156304 Hs.196102	ESTs ESTs	5.60	6.49
	310624	AI341594	,,,,,,,,,,	gb:Human endogenous retrovirus H proteas	4.91	9.09
75	310636	AI814373	Hs.164175	ESTs	1.85	1.71
75	310648 310694	A1347863 A1654370	Hs.156672 Hs.157752	ESTs Homo sapiens mRNA full length insert cDN	0,17 . 5.40	0.69 13.22
	310694	A1472124	Hs.157757	ESTs	4.82	6.27
	310714	AI418446	Hs.157882	ESTs	1.76	3.51
QΛ	310722	AI989803	Hs.157289	ESTs	1.14 8.46	6.85 13.01
80	310756 310764	Al916560 Al376769	Hs.158707 Hs.167172	ESTs ESTs	4.76	7.37
	310848	Al459554	Hs.161286	ESTS	2.84	1.96
	310851	AW291714	Hs.221703	ESTs	1.00	2.32
85	310854	AI421677	Hs.161332	ESTs	6.37 6.07	7.94 9.84
92	310858	AI871000	Hs.161330	ESTs	0.01	3.04

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	310864	Al924558	Hs.161399	ESTs	0.87	0.78
	310875	T47764	Hs.132917	ESTs	1.00	3.63
	310896	AW157731	Hs.270982	ESTs, Moderately similar to ALU7_HUMAN A	7.07	16.68
-	310922	AW195634	Hs.170401	ESTs	1.00	1.00
5	310955	AI560210	Hs.263912	ESTs	10.08 2.18	17.66 3.18
	310957 311000	AW190974 AI521830	Hs.196918 Hs.171050	ESTs	3.06	6.64
	311012	AW298070	Hs.241097	ESTs ESTs	1.23	3.77
	311034	AI564023	Hs.311389	ESTs, Moderately similar to PT0375 natur	2.44	2.09
10	311074	AW290922	Hs.199848	ESTs	6.04	14.19
	311134	A1990849	Hs.196971	ESTs	3.54	6.96
	311174	AW450552	Hs.205457	periaxin	0.65	0.95
	311187	A1638374 A1656040	Hs.224189	ESTs ESTs	2.46 1.10	2.78 2.52
15	311220 311230	A1989808	Hs.196532 Hs.197663	ESTS	1.41	1.75
13	311236	AI653378	Hs.197674	ESTs	2,18	2.11
	311242	AW016812	Hs.200266	ESTs	0.63	5.11
	311258	Al671221	Hs.199887	ESTs	1.00	1.41
20	311277	AW072813	Hs.270868	ESTs, Moderately similar to ALU4_HUMAN A	2.56	1.94
20	311294	AA826425	Hs.291829	ESTS	1.04 1.96	2.69 6.70
	311308 311351	F12664 Al682303	Hs.49000 Hs.201274	ESTs ESTs	4.77	9.38
	311390	AW392997	Hs.202280	ESTs	2.80	6.06
	311405	AW290961	Hs.201815	ESTs	3.80	11.66
25	311409	A1698839		gb:wd31f02.x1 Soares_NFL_T_GBC_S1 Homo s	3.84	6.94
	311420	Al936291	Hs.209867	ESTs	5.30	12.56
	311443	A1791521	Hs.192206	ESTs	4.39 1.00	6.09 1.04
	311467 311479	A1934909 A1933672	Hs.175377 Hs.211399	ESTs ESTs	2.76	5.61
30	311488	R57390	Hs.301064	arfaptin 1	2.50	5.73
-	311495	AW300077	Hs.221358	ESTs .	3.63	6.09
	311511	AW444568	Hs.210303	ESTs	2.00	2.87
	311534	AW130351	Hs.243549	ESTs	0.31	1.33
35	311537	A1805121	Hs.211828	ESTs	3.69 1.73 -	5.85 1.34
33	311543 311551	AI681360 AW449774	Hs.201259 Hs.296380	ESTs POM (POM121 rat homolog) and ZP3 fusion	3.31	6.12
	311557	AIB19230	Hs.211238	interleukin-1 homolog 1	1.00	1.00
	311558	Z44432	Hs.63128	KIAA1292 protein	2.25	3.41
40	311559	AW008271	Hs.265848	similar to rat myomegalin	2.68	5.90
40	311563	AI922143	Hs.211334	ESTs	2.39	3.32
	311586	AI827834	Hs.211227	ESTS	2.47 1.00	3.85 1.00
	311616 311621	AW450675 Al924307	Hs.212709 Hs.213464	ESTs ESTs	4.16	6.74
	311635	AI928456	Hs.213081	ESTs	2.17	3.76
45	311668	AW193674	Hs.240044	ESTs	2.60	3.12
	311672	R11807	Hs.20914	hypothetical protein FLJ23056	2.79	5.18
	311683	AW183738	Hs.232644	ESTs	0.19	0.96
	311700	R49601 AW131785	Hs.171495 Hs.246831	relinoic acid receptor, beta ESTs, Weakly similar to CIKG_HUMAN VOLTA	6.28 5.00	8.83 8.17
50	311714 311735	AW294416	Hs.144687	Homo sapiens cDNA FLJ12981 fis, clone NT	0.96	0.72
-	311743	T99079	Hs.191194	ESTs	1.00	1.95
	311783	AJ682478	Hs.13528	hypothetical protein FLJ14054	0.16	0.77
	311785	A1056769	Hs.133512	ESTs .	1.34	3.97
55	311799	AA780791	Hs.14014	ESTs, Weakly similar to KIAA0973 protein	8.52 3.58	13.32 3.91
55	311819	AW265275 AJ089422	Hs.254325 Hs.131297	ESTs	1.40	1.72
	311823 311877	AA349893	Hs.85339	G protein-coupled receptor 39	0.95	0.91
	311886	AA522738	Hs.132554	ESTs	0.88	0.87
	311896	AW206447		gb:UI-H-BI1-afg-g-02-0-UI.s1 NCI_CGAP_Su	1.66	1.13
60	311910	N28365	Hs.22579	Homo saplens clone CDABP0036 mRNA sequen	1.66	2.30
	311923	T60843	Hs.189679	ESTS	0.42 1.88	2.63 3.02
	311933 311959	AI597963 T67262	Hs.118726 Hs.124733	ESTs ESTs	2.02	2.33
	311960	AW440133	Hs.189690	ESTs	3.87	6.62
65	311967	Al382726	Hs.182434	ESTs	5.80	8.14
٠, .	311975	AA804374	Hs.272203	Homo sapiens cDNA FLJ20843 fis, clone AD	0.98	3.26
	312005	T78450	Hs.13941	ESTs	0.12	1.39
	312028	T78886	Hs.284450	ESTs	3.78 4.11	4.92 7.32
70	312046 312056	A1580018 T83748	Hs.268591 Hs.268594	ESTs ESTs	2.36	3.08
70	312064	AA676713	Hs.191155	ESTs	3.34	5.28
	312088	AW303760	Hs.13685	ESTs	1.60	1.15
	312093	T91809	Hs.121296	ESTs	0.68	0.85
75	312094	Z78390	11- 445456	gb:HSZ78390 Human fetal brain S. Meier-E	3.05	4.48
75	312097	Al352096	Hs.112180	zinc finger protein 148 (pHZ-52)	4.52 2.40	9.70 2.60
	312118 312128	T85332 Al052609	Hs.178294 Hs.17631	ESTs Homo sapiens cDNA FLJ20118 fis, clone CO	2.39	3.53
	312147	T89855	Hs.195648	ESTs	0.67	1.03
••	312175	AA953383	Hs.127554	ESTs	5.85	10.60
80	312179	AI052572	Hs-269864	ESTs	2.41	3.32
	312201	Al928365	Hs.91139	solute carrier family 1 (neuronal/epithe	0.24	0.89 4.55
	312207 312220	H90213 N74613	Hs.191330	ESTs gb:za55a07.s1 Soares fetal liver spleen	2.20 4.28	4.55 11.13
	312252	Al128388	Hs.143655	ESTs	1.64	1.57
85	312304	AA491949	Hs.269392	ESTs	0.12	2.47

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	313926	AW473830	Hs.171442	ESTs	3.40	4.11
	313948	AW452823	Hs.135268	ESTs	5.77	9.15
	313978	AI870175	Hs.13957	ESTs .	0.46	0.75 6.40
5	313983 314035	AI829133 AA164199	Hs.226780 Hs.270152	ESTs	4.10 5.88	7.90
,	314037	AW300048	Hs.275272	ESTs ESTs	1.00	3.79
	314040	AA166970	Hs.118748	ESTs	7.60	11.33
	314067	AW293538	Hs.51743	KIAA1340 protein	1.86	1.21
	314103	AI028477	Hs.132775	ESTs	2.90	5.29
10	314107	AA806113	Hs.189025	ESTs	2.00	1.66
	314113	AA218986	Hs.118854	ESTs	0.91	4.17
	314124	AW118745	Hs.9460	Homo sapiens mRNA; cDNA DKFZp547C244 (fr	2.53	3.32
	314126	AA226431		gb:nc18b12.s1 NCI_CGAP_Pr1 Homo sapiens	3.13	5.08
15	314128	AA935633	Hs.194628	ESTs	2.90	6.35
13	314151	AA236163	Hs.202430	ESTs	4.15 3.44	6.45 4.65
	314184 314192	AW081795 AW290975	Hs.233465 Hs.118923	ESTs ESTs	1.00	1.23
	314244	AL036450	Hs.103238	ESTs	2.88	3.67
	314253	AA278679	Hs.189510	ESTs	4.98	7.16
20	314262	AW086215	Hs.246096	ESTs	0.38	1.94
	314320	AA811598	Hs.275809	ESTs	3.34	5.66
	314332	AL037551	Hs.95612	ESTs	2.85	2.09
	314335	AA287443	Hs.142570	Homo saplens clone 24629 mRNA sequence	4.35	4.78
05	314340	AW304350	Hs.130879	ESTs, Moderately similar to putative p15	0.77	0.86
25	314351	AA292275	Hs.193746	ESTs	3.07	3.77
	314376	AJ628633	Hs.324679	ESTs	4.10	6.11
	314443	AA827125	Hs.192043	ESTs	6.20	13.67 2.49
	314458 314466	A1217440 AA767818	Hs.143873 Hs.122707	ESTs ESTs	0.58 2.53	2.49
30	314478	AI521173		DEAD-box protein	3.94	5.65
50	314482		Hs.134182	ESTs	1.30	1.44
	314506	AA833655	Hs.206868	Homo sapiens cDNA FLJ14056 fis, clone HE	3.28	3.47
	314519	R42554	Hs.210862	T-box, brain, 1	3.12	6.16
	314529	AL046412	Hs.202151	ESTs	3.43	6.87
35	314546	AW007211	Hs.16131	hypothetical protein FLJ12876	1.38	1.00
	314562	AI564127	Hs.143493	ESTs	2.29	5.27
	314579	AW197442	Hs.116998	ESTs	3.87	5.75
	314580	AW451832	Hs.255938	ESTs, Moderately similar to KIAA1200 pro	0.10	0.71
40	314585	AA918474	Hs.216363	ESTS Home conjugate aDNA EL 110570 for clone NT	1.08 1.00	1.40 1.00
40	314589 314592	AW384790 AA435761	Hs.153408 Hs.192148	Homo sapiens cDNA FLJ10570 fis, clone NT ESTs	0.90	2.60
	314603	AA418024	Hs.270670	ESTs	4.56	6.29
	314604	AA946582	Hs.8700	deleted in liver cancer 1	3.42	3.92
	314606	AA418241	Hs.188767	ESTs	2.97	4.55
45	314648	AA878419		gb:EST391378 MAGE resequences, MAGP Homo1	.42	1.36
	314699	AI038719	Hs.132801	ESTs	3.66	4.97
	314701	Al754634	Hs.131987	ESTs	0.03	0.90
	314710	AI669131	Hs.290989	EST	3.40	7.52
50	314750	AI095005	Hs.135174	ESTs	2.80	6.54
50	314767	AW135412	Hs.164002	ESTs	3.20	4.26
	314801	AA481027	Hs.109045	hypothetical protein FLJ10498	1.00 0.91	1.00 0.99
	314817 314835	Al694139 Al281370	Hs.192855 Hs.76064	ESTs ribosomal protein L27a	5.75	7.44
	314852	Al903735	113.10004	gb:MR-BT035-200199-031 BT035 Homo saplen	1.68	4.34
55	314853	AA729232	Hs.153279	ESTs	0.60	1.85
	314940	AW452768	Hs.162045	ESTs	10.10	16.20
	314941	AA515902	Hs.130650	ESTs	0.31	1.02
	314943	Al476797	Hs.184572	cell division cycle 2, G1 to S and G2 to	2.18	0.37
C O	314955	AA521382	Hs.192534	ESTs	2.59	3.90
60 ⋅ ′	314973	AW273128	Hs.300268	ESTs	1.05	1.25
	315004	AA527941	Hs.325351	EST	5.64	13.63
	315006	Al538613	Hs.298241	Transmembrane protease, serine 3	0.52	1.78
	315033	A1493046	Hs.146133	ESTs ESTs	2.46 0.34	1.00 1.33
65	315035 315056	A1569476 A1202703	Hs.177135 Hs.152414	ESTs	2.10	2.64
05	315069	AI821517	Hs.105866	ESTs	1.00	1.30
	315071	AA552690	Hs.152423	Homo sapiens cDNA: FLJ21274 fis, clone C	1.78	1.00
	315073	AW452948	Hs.257631	ESTs	1.17	1.52
	315078	AA568548	Hs.190616	ESTs	3.00	3.79
70	315080	AA744550	Hs.136345	ESTs	1.00	1.00
	315120	AA564991	Hs.269477	ESTs	0.64	1.44
	315175	AJ025842	Hs.152530	ESTs	0.61	1.91
	315193	AI241331	Hs.131765	ESTs	1.06	0.97
75	315196	AA972756	Hs.44898	Homo sapiens clone TCCCTA00151 mRNA sequ	0.48	1.96
75	315200	A1808235	Hs.307686	EST	3.76 5.37	9.40
	315254	A1474433	Hs.179556	ESTs hypothetical protein FLJ10493	5.37 1.00	9.36 1.30
	315353 315397	AW452608 AA218940	Hs.279610 Hs.137516	fidgetin-like 1	3.38	2.24
	315403	AW362980	Hs.163924	ESTs	2.04	5.23
80	315431	AA622104	Hs.184838	ESTS	2.36	8.04
	315454	Al239473		gb:qh36f02.x1 Soares_NFL_T_GBC_S1 Homo s	3.46	7.64
	315455	AW393391	Hs.156919	ESTs	3.78	5.76
	315473	AI681671	Hs.312671	ESTs, Moderately similar to OVCA1	0.89	2.15
05	315483	AW512763	Hs.222024	transcription factor BMAL2	2.32	1.96
85	315526	Al193048	Hs.128685	ESTs	1.67	1.78

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	315530	Af200852	Hs.127780	ESTs	1.05	1.01
	315541	Al168233	Hs.123159	sperm associated antigen 4	0.85	0.56
	315552	AW445034	Hs.256578	ESTs	1.00	2.22
_	315562	AA737415	Hs.152826	ESTs	2.66	2.48
5	315577	AW513545	Hs.17283	hypothetical protein FLJ 10890	2.20 1.00	2.25 1.04
	315587	Al268399 AW072387	Hs.140489 Hs.158258	ESTs Homo santens mRNA; cDNA DKFZp434B1272 (f	0.14	1.05
	315589 315623	AA364078	Hs.258189	ESTs	7.44	12.56
_	315634	AA837085	Hs.220585	ESTs	0.50	1.40
10	315668	AA912347	Hs.136585	ESTs	0.43	1.22
	315677	Al932662	Hs.164073	ESTs	0.60	1.39
	315706	AW440742	Hs.155556	hypothetical protein FLJ20202	2.18 2.88	3.77 2.63
	315707	Al418055 H25899	Hs.161160 Hs.201591	ESTs ESTs	0.11	0.60
15	315730 315745	Al821759	Hs.191856	ESTs	3.50	7.25
10	315791	AA678177	110.101000	gb:zi15a05.s1 Soares_fetal_liver_spleen_	1.78	2.63
	315801	AA827752	Hs.266134	ESTs	4.31	6.23
	315820	Al652022	Hs.258785	ESTs	2.35	3.01
20	315878	AA683336	Hs.189046	ESTs	2.12 1.03	2.64 1.97
20	315905 315923	AI821911 AI052789	Hs.209452 Hs.133263	ESTs ESTs	2.63	5.06
	315954	AW276810	Hs.254859	ESTs, Moderately similar to ALU5_HUMAN A	1,21	0.85
	315978	AA830893	Hs.119769	ESTs	3.09	3.41
	316001	AJ248584	Hs.190745	Homo sapiens cDNA: FLJ21326 fis, clone C	2.20	6.82
25	316011	AW516953	Hs.201372	ESTs	0.35	1.63
	316012	AA764950	Hs.119898	ESTs	6.56 5.69	8.13 10.69
	316040 316048	Al983409 Al720759	Hs.189226 Hs.224971	ESTs ESTs	2.84	10.45
	316076	AW297895	Hs.116424	ESTs	0.30	1.05
30	316124	Al308862	Hs.167028	ESTs	1.00	1.43
-	316151	AI806016	Hs.156520	ESTs	5.80	9.03
	316187	AW518299	Hs.192253	ESTs	1.20	3.96
	316204	AA731509	Hs.120257	ESTs	4.92 1.48	6.94 1.60
35	316232 316275	AW297853 Al671041	Hs.251203 Hs.292611	ESTs ESTs, Moderately similar to ALU1_HUMAN A	5.86	12.14
33	316291	AW375974	Hs.156704	ESTs	2.73	2.69
	316303	AA740994	Hs.209609	ESTs	1.53	1.26
	316344	AA744518	Hs.120610	ESTs	3.66	8.34
40	316346	Al028478	Hs.157447	ESTs	3.51	6.69
40	316365	AI627845	Hs.210776	ESTs	2.50 1.16	4.33 2.16
	316380 316470	A1393378 AA809902	Hs.164496 Hs.243813	ESTs ESTs	5.40	10.34
	316509	AA767310	Hs.291766	ESTs	2.46	2.89
	316514	AA768037	Hs.291671	ESTs	4.70	6.04
45	316519	Al929097		gb:od10c11.s1 NCI_CGAP_GCB1 Homo sapiens	4.41	9.70
	316609	AW292520	Hs.122082	ESTs	1.00	2.89 3.72
	316633	Al125586 AW172316	Hs.127955 Hs.252961	ESTs ESTs, Weakly similar to ALU1_HUMAN ALU S	2.61 3.46	4.64
	316700 316711	AI743721	Hs.285316	ESTs, Moderately similar to ALU7_HUMAN A	4.45	6.95
50	316713	AI090671	Hs.134807	hypothetical protein FLJ12057	0.30	2.40
	316715	Al440266	Hs.170673	ESTs, Weakly similar to AF126780 1 retin	0.20	1.45
	316787	AW369770	Hs.130351	ESTs	4.05	5.53
	316809	AA825839	Hs.202238	ESTs	2.25 1.00	3.82 1.32
55	.316811 316812	AA922060 AW135045	Hs.132471 Hs.232001	ESTs ESTs	3.28	4.70
55	316818	AA827176	Hs.124316	ESTs	0.67	1.81-
	316824	AA837416	Hs.124299	ESTs	3.53	6.00
	316827	Al380429	Hs.172445	ESTs	0.72	1.56
<i>c</i> ο .	316891	AW298119	Hs.202536	ESTs	1.64 1.45	2.97 1.08
60	316951	AA134365 AA860172	Hs.57548 Hs.132406	ESTs ESTs	1.00	1.53
	316970 316971	AA860212	Hs.170991	ESTs	1.08	1.96
	316990	AA861611	Hs.130643	ESTs	5.44	10.04
	317001	AI627917	Hs.233694	hypothetical protein FLJ11350	3.56	4.37
65	317008	AW051597	Hs.143707	ESTs	0.69	1.37
	317051	AA873253	Hs.126233	ESTs ESTs	6.18 1.87	12.72 2.66
	317128 317129	AA971374 H12523	Hs.125674 Hs.78521	ESTs Homo sapiens cDNA: FLJ21193 fis, clone C	4.12	6.64
	317125	AW341567	Hs.125710	ESTs	2.82	5.12
70	317196	Al348258	Hs.153412	ESTs	1.98	2.51
	317212	A1866468	Hs.148294	ESTs	1.86	2.83
	317223	AW297920	Hs.130054	ESTs	0.83	1.57
	317224	D56760	Hs.93029	sparc/osteonectin, cwcv and kazal-like d	2.74 1.00	0.86 1.00
75	317266 317282	AA906289 Al807444	Hs:203614 Hs:176101	ESTs ESTs	2.60	4.21
	317285	AW370882	Hs.222080	ESTs	1.96	3.49
	317302	AA908709	Hs.135564	ESTs	7.16	8.32
	317304	AW449899	Hs.130184	ESTs	1.38	2.28
90	317320	AA927151	Hs.130452	ESTs	3.58	8.13 4.92
80	317413	AW341701	Hs.126622	ESTs ESTs	2.08 3.06	4.92 4.79
	317417 317452	AA918420 AA972965	Hs.145378 Hs.135568	EST8	4.22	9.21
	317519	A1859695	Hs.126860	ESTs	1.88	4.15
ò۶	317521	A182433B	Hs.126891	ESTs	3.12	4.55
85	317529	A1916517	Hs.126865	EST8	2.73	3.34

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	317570	AJ733361	Hs.127122	ESTs	1.00	2.43
	317571	AA938663	Hs.199828	ESTs	5.20	11.95
	317598	AW206035	Hs.192123	ESTs	0.33	1.56
5	317627 317650	Al346110 Al733310	Hs.132553 Hs.127346	ESTs ESTs	1.50 0.48	1.39 1.46
,	317659	AA961216	Hs.127785	ESTs	4.18	7.14
	317674	AW294909	Hs.132208	ESTs	2.92	3.20
	317686	AA969051	Hs.187319	ESTs	1.00	1.01 9.59
10	317692 317701	Al307659 Al674774	Hs.174794 Hs.128014	ESTs ESTs	5.33 1.00	1.00
10	317711	Al733015	Hs.272189	ESTs	5.13	7.81
	317722	A1733373	Hs.128119	ESTs	2.50	6.03
	317756	AA973667	Hs.128320	EST8	1.59	1.30
15	317777 317799	Al143525 Al498273	Hs.47313 Hs.128808	KIAA0258 gene product ESTs	1.00 1.78	2.48 2.11
13	317803	AA983251	Hs.128899	ESTs	0.80	1.06
	317821	AJ368158	Hs.70983	PTPL1-associated RhoGAP 1	0.17	0.68
	317848	AI820575	Hs.129086	Homo saptens cDNA FLJ12007 fis, clone HE	5.30	8.16
20	317850	N29974	Hs.152982 Hs.129119	hypothetical protein FLJ13117 ESTs	1.30 2.18	2.28 5.93
20	317861 317865	AW341064 Al298794	Hs.129130	ESTS	4.48	8.20
	317869	AW295184	Hs.129142	deoxyribonuclease II beta	0.44	0.99
	317881	AI827248	Hs.224398	Homo sapiens cDNA FLJ11469 fis, clone HE	4.06	2.23
25	317890 317899	Al915599 Al952430	Hs.129225 Hs.150614	ESTs ESTs, Weakly similar to ALU4_HUMAN ALU S	4.68 3.14	7.48 3.37
25	317986	A1005163	Hs.201378	ESTs, Weakly similar to T12545 hypotheti	0.28	1.66
	318001	AW235697	Hs.130980	ESTs	5.12	9.97
	318016	AI016694	Hs.256921	ESTs	1.86	4.50
30	318023	AW243058	Hs.131155	ESTs ESTs	2.92 3.92	5.22 6.37
30	318054 318068	AW449270 Al024540	Hs.232140 Hs.131574	ESTS	1.21	1.27
	318117	AI208304	Hs.250114	ESTs	0.86	1.17
	318187	A1792585	Hs.133272	ESTs, Weakly similar to ALUC_HUMAN IIII	5.90	6.98
35	318223	A1077540	Hs.134090	ESTs	1.05 3.10	0.90 2.40
33	318240 318255	Al085377 Al082692	Hs.143610 Hs.134662	ESTs ESTs	0.02	1.05
	318266	AI554341	Hs.271443	ESTs	6.12	10.55
	318330	A1093840	Hs.143758	ESTs	4.98	7.90
40	318369	Al493501	Hs.170974	ESTs ESTs	2.46 0.77	5.62 0.45
40	318428 318458	A1949409 A1149783	Hs.194591 Hs.158438	ESTS	3.54	4.92
	318467	Al151395	Hs.144834	ESTs	4.56	5.62
	318473	A1939339	Hs.146883	ESTs	2.08	4.05
45	318476	AI693927	Hs.265165	ESTs ESTs	4.22 1.47	8.07 1.05
45	318487 318488	Al167877 Al217431	Hs.143716 Hs.144709	ESTs ESTs	1.40	4.14
		- T26477	Hs.22883	ESTs, Weakly similar to ALU8_HUMAN ALU S	1.84	1.90
	318499	T25451		gb:PTHI188 HTCDL1 Homo sapiens cDNA 5/3	2.58	5.20
50	318537 318538	AA377908 N28625	Hs.13254 Hs.74034	ESTs Homo sapiens clone 24651 mRNA sequence	3.26 0.35	4.18 1.07
50	318547	R20578	Hs.90431	ESTs	3.22	4.60
	318552	R18364	Hs.90363	ESTs	4.87	9.06
	318575	R55102	Hs.107761	ESTs, Wealdy similar to unnamed protein	1.91	1.98
55	318580	T34571 AA779704	Hs.49007	poly(A) polymerase alpha Homo sapiens cDNA FLJ12136 fis, clone MA	2.74 0.85	6,22 2.46
55	318587 318596	AI470235	Hs.168830 Hs.172698	EST	4.88	4.93
	318622	T48325	Hs.237658	apolipoprotein A-II	4.80	12.51
	318629	N25163	Hs.8861	ESTs	0.39	1.04
60	318637 318648	AA243539 T77141	Hs.9196 Hs.184411	hypothetical protein albumin	1.72 6.27	3.57 9.91
00	318650	AA393302	Hs.176626	hypothetical protein EDAG-1	3.96	8.84
	318671	AA188823	Hs.299254	Homo sapiens cDNA: FLJ23597 fis, clone L	1.53	0.81
	318679	T58115	Hs.10336	ESTs	1.00	2.19
65	318711 318725	Al936475 Al962487	Hs.101282 Hs.242990	Homo sapiens cDNA: FLJ21238 fis, clone C ESTs	3.05 1.08	3.18 2.46
05	318728	Z30201	Hs.291289	ESTs, Weakly similar to ALU1_HUMAN ALU S	0.77	1.33
	318740	NM_002543		oxidised low density lipoprotein (lectin	0.25	1.49
	318776	R24963	Hs.23766	ESTs	1.00	3.01
70	318784 318816	H00148 F07873	Hs.5181 Hs.21273	proliferation-associated 2G4, 38kD ESTs	2.70 3.90	3.86 7.13
70	318865	H10818	113.21213	gb:ym04f10.r1 Soares infant brain 1NIB H	2.25	3.56
	318879	R56332	Hs.18268	adenylate kinase 5	1.78	5.00
	318881	Z43224	Hs.124952	ESTs	4.79	14.13
75	318894	F08138 AW368520	Hs.7387 Hs.301528	DKFZP564B116 protein L-kynurenine/alpha-aminoadipate aminotra	5.31 1.03	7.00 0.91
15	318901 318925	Z43577	Hs.21470	ESTs	2.23	3.80
	318936	Al219221	Hs.308298	ESTs	1.86	7.16
	318982	Z44140	Hs.269622	ESTs	5.84	9.79
80	318986	Z44186	Hs.169161	ESTs, Highly similar to MAON_HUMAN NADP- ESTs, Weakly similar to weak similarity	1.00 3.38	1.00 6.11
50	319041 319103	Z44720 H05896	Hs.98365 Hs.4993	KIAA1313 protein	1.00	1.07
	319170	R13678	Hs.285306	putative selenocysteine tyase	3.79	5.03
	319196	F07953	Hs.16085	putative G-protein coupled receptor	1.00	2.98
85	319199	F07361	Hs.13306	ESTs ESTs	3.53 5.87	5.66 7.26
UJ.	319242	F11472	Hs.12839	ESTs	3.01	7.20

	W	O 02/080	5443			
	319263	T65331	Hs.81360 .	Homo sapiens cDNA: FLJ21927 fis, clone H	1.81	1.57
	319267	F11802	Hs.6818	ESTs	1.10	4.72
	319270	R13474	Hs.290263	ESTs	4.80	10.40
_	319279	T65094	Hs.12677	CGI-147 protein	1.50	211
5	319282	AA461358	Hs.12876	ESTs	1.00	1.00
_	319289	W07304	Hs.79059	transforming growth factor, beta recepto	0.18	0.68
	319291	W86578	Hs.285243	hypothetical protein FLJ22029	0.26	0.62
	319293	F12119	Hs.12583	ESTs	3.13	4.50
	319312	Z45481	1.0.12000	gb:HSC2QE041 normalized infant brain cDN	1.10	1.00
10	319370	H54254	Hs.325823	ESTs, Moderately similar to ALU5_HUMAN A	0.16	0.73
10	319391	R06304	Hs.13911	ESTs	1.26	2.43
	319396	H67130	Hs.301743	ESTs	0.70	0.76
	319398	AA359754	Hs.191196	ESTs	2.45	3.59
	319407	R05329	NS.131130	gb:ye91b04.r1 Soares fetal liver spleen	2.00	3.54
15				gb:yd39f07.r1 Soares fetal liver spleen	4.28	8.81
13	319425	T82930	U- 404400	* . <u>*</u>	6.15	
	319433	R06050	Hs.191198	ESTS		14.13
	319437	AA282420	Hs.111991	ESTs, Weakly similar to Y48A5A.1 [C.eleg	3.26	5.68
	319466	AI809937	Hs.116417	ESTs	1.76	5.65
20	319471	R06546	Hs.19717	ESTs	4.29	4.84
20	319480	R06933	Hs.184221	ESTs	1.00	1.00
	319484	T91772		gb:yd52a10.s1 Soares felal liver spleen	2.81	4.88
	319486	A1382429	Hs.250799	ESTs	2.08	2.82
	319508	T99898	Hs.270104	ESTs, Moderately similar to ALU8_HUMAN A	2.80	4.39
	319523	T69499	Hs.191184	ESTs	1.55	3.25
25	319545	R83716	Hs.14355	Homo sapiens cDNA FLJ13207 fis, clone NT	1.65	1.19
	319546	R09692		gb:yf23b12.r1 Soares fetal liver spleen	5.11	8.54
	319552	AA096106	Hs.20403	ESTs	1.89	3.36
	319582	T82998	Hs.250154	hypothetical protein FLJ12973	3.48	4.82
	319586	D78808	Hs.283683	chromosome 8 open reading frame 4	0.26	0.82
30	319604	R11679	Hs.297753	vimentin	1.68	3.41
50	319609	AW247514	Hs.12293	hypothetical protein FLJ21103	3.06	4.24
			115.12235	gb:ym19c10.r1 Soares infant brain 1NIB H	2.76	4.24
	319611	H14957	Hs.173515		2.51	3.55
	319653	AA770183		uncharacterized hypothalamus protein HT0	5.32	7.68
35	319657	R19897	Hs.106604	ESTs		
22	319658	R13432	Hs.167481	syntrophin, gamma 1	3.35	5.00
	319661	H08035	Hs.21398	ESTs, Moderately similar to A Chain A, H	5.18	12.55
	319662	H06382	Hs.21400	ESTs	1.58	1.56
	319708	R15372	Hs.22664	ESTs	1.00	1.22
40	319742	T77668	Hs.21162	ESTs	2.48	3.13
40	319748	R18178	Hs.295866	Homo sapiens mRNA; cDNA DKFZp434N1923 (f	3.02	4.85
	319772	R76633	Hs.22646	ESTs	4.36	11.61
	319788	AA321932	Hs.117414	KIAA1320 protein	2.56	3.68
	319805	R92857	Hs.271350	likely ortholog of mouse polydom	4.63	6.56
	319812	N74880	Hs.264330	N-acylsphingosine amidohydrolase (acid c	0.63	1.32
45	319834	AA071267		gb:zm61g01.r1 Stratagene fibroblast (937	0.30	0.94
	319878	T78517	Hs.13941	ESTs	3.99	6.44
	319882	AA258981	Hs.291392	ESTs ·	5.09	7.36
	319912	T77559	Hs.94109	Homo sapiens cDNA FLJ13634 fis, clone PL	3.24	3.21
•	319935	H79460	Hs.271722	ESTs, Weakly similar to ALU1_HUMAN ALU S	4.40	9.42
50	319944	T79248	Hs.133510	ESTs	3.31	5.39
•	319947	AA160967	Hs.14479	Homo sapiens cDNA FLJ14199 fis, clone NT	2.90	4.95
	319962	H06350	Hs.135056	Human DNA sequence from clone RP5-850E9	1.81	1.57
	320007	AA336314	113.100000	gb:EST40943 Endometrial tumor Homo saple	3.42	6.29
	320018	T83263		gb:yd40h09.r1 Soares fetal liver spleen	2.77	5.14
55	320030	H63789	Hs.296288	ESTs, Weakly similar to KIAA0638 protein	4.10	6.69
<i>J J</i>	320030				3.27	3.27
		A1699772	Hs.292664	ESTs, Weakly similar to A46010 X-linked		
	320040	AA233671	Hs.87164	hypothetical protein FLJ14001	1.81	1.64
	320047	T86564	Hs.302256	EST	3.38	7.36
60	320063	AA074108	Hs.120844	FOXJ2 forkhead factor	5.90	16.73
OU	320096	H58138	Hs.117915	ESTs	2.08	4.47
	320099	AW411307	Hs.114311	CDC45 (cell division cycle 45, S.cerevis	1.00	1.00
	320112	T92107	Hs.188489	ESTs	2.27	2.06
	320140	H94179	Hs.119023	SMC2 (structural maintenance of chromoso	1.00	1.00
CE	320188	AW419200	Hs.172318	ESTs ·	1.26	1.00
65	320193	AA831259	Hs.17132	ESTs	2.58	6.23
	320195	R62203	Hs.24321	Homo sapiens cDNA FLJ12028 fis, clone HE	2.85	4.53
	320199	R78659	Hs.29792	ESTs	0.40	0.94
	320203	AL049227	Hs.124776	Homo sapiens mRNA; cDNA DKFZp564N1116 (f	0.84	1.18
	320219	AA327564	Hs.127011	tubulointerstitial nephritis antigen	1.00	1.17
70 ·	320220	AF054910	Hs.127111	tektin 2 (testicular)	0.18	1.09
	320225	AF058989	Hs.128231	G antigen, family B, 1 (prostate associa	5.26	13.75
	320231	H03139	Hs.24683	ESTs	1.59	1.93
•	320260	NM_003608		G protein-coupled receptor 65	1.38	4.56
	320267	AL049337	Hs.132571	Homo saplens mRNA; cDNA DKFZp564P016 (fr	1.00	1.92
75	320268	H06019	Hs.151293	Homo sapiens cDNA FLJ10664 fis, clone NT	5.58	5.70
	320322	AF077374	Hs.139322	small proline-rich protein 3	1.41	1.01
	320325	AI167978	Hs.139851	caveolin 2	0.05	0.67
	320323	AF026004	Hs.141660	chloride channel 2	2.17	1.26
	320339	H10807	Hs.281434	Homo sapiens cDNA FLJ14028 fis, clone HE	1.81	2.32
80	320388		. Hs.31286	ESTs	1.00	3.22
-	320308			Homo sapiens clone FLC0578 PRO2852 mRNA,	1.41	1.36
		R22291	Hs.23368		2.31	3.61
	320413	AA203711	Hs.173269	ESTS		20.78
	320432	R62786	Hs.124136	ESTs	11.25	
85	320438	AA253352	Hs.293663	ESTS	2.22	3.49
O)	320438	W24548	Hs.5669	EST8	3.53	8.14

320448 AA09443		W	O 02/08	6443			
200451 725944 14.180777					v-ves-1 Yamaquchi sarcoma viral related	1.42	3.46
2049 A.94345 h. 1.98267 h. 1.9826							0.81
30498 Pizzlis Haboro Fibra 20221 A 20222 A 202						0.65	1.18
20251 N3164 Hs.24743 (2025) AV074205 Hs.11314 ESTs 3.66 1.04 20255 AV074205 Hs.132724 Hs.13724 Hs.13724 Hs.13724 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 20255 AF05417 Hs.14570 hypothetical protein FLJ22530 1.28 1.00 20255 AF05417 Hs.14570 hypothetical protein FLJ22530 1.28 1.00 20255 AF05417 Hs.14570 hypothetical protein FLJ22530 1.28 1.00 20255 AF05427 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 20255 AF05427 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 20255 AF05427 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 2.00 2025 AF05427 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 20255 AF05427 Hs.15995 pelphylglyche alpha-emidating monoxyg 1.22 1.00 2.00 2.00 2.00 2.00 2.00 2.00	_	320499	R32555		Homo sapiens cDNA FLJ12028 fis, clone HE	3.44	7.15
2002579 AV347205 Hs.111314 ESTs	5		AB007978	Hs.158278	KIAA0509 protein		
200527 ASSASS AASS1724 htts. 14570 200526 AF054771 htts. 14570 200526 AF054771 htts. 14570 200526 AF055209 htts. 15930 https://doi.org/10.10016/10.10016/20.20053 AF055209 htts. 15930 https://doi.org/10.10016/20.20053 AF055209 htts. 15930 https://doi.org/10.10016/20.20053 AASSASS htts. 15930 ht							
20255 APAS1772 Hs.157272							
2005 AP054077 Hs.14570 2005 AP055079 Hs.15335 hs.15355 hs							
320654 AF056209 htt.159386 peptlyfyglybne alpha-emidaling morouxyg 1.22 0.041	1Λ						
200535	10						
200353 PA.159 H. 80508 Small nuclear Thorouseporcelin polypept 1.00 6.25							
20838 A84288 H. 17335 hypothetical protein FLU3182							
15 20064 MASS21 H.20559 Homo sapiner mRNA for KIAA1708 protein, 1.00 1.53 20064 AA402269 H.511334 H.510510 H							
200664	15						
20066						0.14	0.79
20 320638 R52921		320664	Al904216	Hs.91251			
20 320839 AJ34699 Hs.171929 ESTs Westly similar to ASA849 collagen 1.27 1.02 20056 AV135016 Hs.172780 gby, 40451913 200771 J8792366 Hs.172780 gby, 40451913 Hs.172780 gby, 4045191 Hs.172780 gby, 40		320676	AA132650	Hs.300511	ESTs		
3209114 AM45591 hs.17176 syppolda10.r1 Soares fotal liver spleen 1.06 0.85	20						
20071 A A A A A Sept	20						
257 20097				Hs.172780			
25 32077 Al A281933 Hs. 191276 Js. 19679 Js. 1				U- 10112E			
25 320794 AA21993 Hs. 91265 ESTs							
320822	25						
320824 AF120274 hs.194689 arlamin claudin 14 .106 1.11							
30830						1.16	1.11
300 320849 060031 Hs.34771 ESTs					claudin 14	1.06	1.75
320853 A473796 Hs. 135904 ESTs Log 1.00		320843	AA317372	Hs.34744	Homo sapiens mRNA; cDNA DKFZp547C136 (fr		
320985	30	320849	D60031	Hs.34771	ESTs		
320927 A920786 Hs. 199338 Hs. 199338 Hs. 201707 A920787 A920787 A9207833 Hs. 201823 Core histone macroH2A2.2 1.67 2.18 2.1045 W38483 Hs. 290350 ESTs 2.69 4.25 4.55 3.2046 A920784 Hs. 209055 ESTs 2.69 4.25 4.55 3.2062 A920784 Hs. 204101 A920784 Hs. 204101 A920784 Hs. 204101 A920784 A920884 Hs. 204101 A920784 A920884 Hs. 204101 A920784 A920884 Hs. 204101 A92088 A920884 A92088							
320927 A12057786 Hs.213923 STR Core histone macroH2A2_2 16.7 2.18 2.18 2.21 2.14 2.18 2.19 2.10 2.18 2.10 2.18 2.10 2.18 2.10 2.18 2.10 2.18 2.10 2.18 2.10 2.18 2.10							
1.592037 1.592033 1.5.92							
320997 H22544 321045 W88483 Hs.293650 ESTs 2.25 4.55 4.55 321046 H27794 Hs.269055 ESTs 2.69 4.25 4.55 321052 AU872284 Hs.2640770 Muclear cap binding protein subunit 2, 2 2.14 2.56 2.25 4.55 321052 AU872284 Hs.126454 ESTs 2.25 4.55 321052 AU87284 Hs.241413 Momo sapiens clone 24941 mRNA sequence 4.79 7.41 4.4750 Hs.125494 ESTs 4.366635 Au872894 Hs.125494 ESTs 4.366635 Au872894 Hs.125494 ESTs 4.57 Au873833 Hs.298351 Au872899 321170 Au873834 Hs.29598 Au872894 Hs.226469 Hs.1174 Au87480 Hs.125494 Hs.226469 Au87480	25						
321045 W88483 Hs.293650 ESTs 2.25 4.55	33			ns.32023			
321046 H27794 Hs.269055 ESTs 2.69 4.25				Hs 293650			
August Hs.24977 String Hs.24977 String August Hs.24978 Hs.241411 Homo sapiens mRNA full length insert cDN 2.76 5.20 5.							
321062 R87955						2.14	2.56
321067	40	321059	A1092824	Hs.126465	ESTs		
\$\frac{321102}{321130} \$A018306 \$\frac{1}{321142} \$A1817933 \$\frac{1}{329351} \$A5PL protein \$8.73 \$15.35 \$321155 \$A336635 \$\frac{1}{321170} \$A27170 \$A5712 \$A316635 \$A59598 \$\frac{1}{321170} \$A3742 \$\frac{1}{321170} \$A35742 \$\frac{1}{321170} \$A5742 \$\frac{1}{321235} \$A003073 \$\frac{1}{321235} \$A003073 \$\frac{1}{321240} \$A1080073 \$\frac{1}{321235} \$A00371941 \$\frac{1}{321235} \$A00371941 \$\frac{1}{34137050} \$\frac{1}{321270} \$\frac{1}{3321270} \$\		321062	R87955	Hs.241411			
45 321142 ABIT933 Hs.298361 Hs.29837 ASPL protein MGC5338 3.04 5.03 321155 AA36635 Hs.99598 hypothetical protein MGC5338 3.04 5.03 321156 AA700289 321170 N53742 Hs.172982 ESTs 221 4.46 321199 AW382512 hs.298381 hypothetical protein MGC5338 3.04 5.03 321199 AW382512 hs.272982 ESTs 221 4.46 321199 AW382512 hs.251414 htmo sapiens cDNA FL.172417 fis, clone MA 4.00 7.32 321236 AW371941 hs.18192 htmo sapiens cDNA FL.172417 fis, clone MA 4.00 7.32 321244 AF058654 hs.26149 htmo sapiens cDNA FL.172417 fis, clone MA 4.00 7.32 321244 AF058654 hs.2628 hs.2628 klaA1161 protein firmulation from the firmu				Hs.241438			
ASP							
321158 AA33635 Hs.99598 hypothetical protein MGC5338 3.04 5.03 321158 AA700289 gb.yu76611.r1 Soares fetal liver spleen 4.62 8.39 21170 N53742 Hs.172982 ESTs 2.21 4.46 339 211206 H54178 Hs.226469 Homo sapiens cDNA FLJ12417 fis, clone MA 4.00 7.32 12226 AL080073 Hs.251414 Homo sapiens mRNA; cDNA DKFZp56481462 (f 4.17 4.63 321236 AW371941 Hs.18192 gb.yv76c05.s1 Soares fetal liver spleen 4.00 7.32 12270 R35560 gb.yu76c05.s1 Soares fetal liver spleen 3.80 5.26 4.32 1317 Al937060 Hs.6298 kiAA161 protein 1.81 1.65 321318 AB033010 Hs.300646 KiAA protein (similar to mouse paladin) 0.44 0.93 121325 AB033100 Hs.300646 KiAA protein (similar to mouse paladin) 0.44 0.93 121325 AB033101 Hs.161075 ESTs 3.1402 A368667 Hs.132743 ESTs 1.13 0.97 321430 U05890 321467 X13075 45.321489 AK251879 Hs.255436 ESTs 2.28 2.54 A51498 AW295517 Hs.255436 ESTs 2.21498 AW295517 Hs.255436 ESTs 2.21490 A036869 Hs.259309 321510 AA703650 Hs.258980 ESTs 2.21491 A036869 Hs.258748 ESTs 2.214 3.94 A036850 Hs.258748 ESTs 2.21491 A036869 Hs.258748 ESTs 2.21491 A036869 Hs.258748 ESTs 2.21491 A036869 Hs.258748 ESTs 2.214 3.94 Hs.258748 Hs.25873 Hs.256514 hypothetical protein FLJ11342 4.89 7.82 321510 AA703650 Hs.258748 ESTs 2.214 3.94 Hs.2660 A295430 Hs.268514 hypothetical protein FLJ11340 A03964 Hs.268514 hypothetical protein FLJ21560 Hs.26869 Hs.26898 Hs.268518 hypothetical protein FLJ21560 Final N204HR Homo appliens CDNA FLJ12830 fis, clone NT 2.82 3.288 3.21688 Hs/7646 Hs.261051 Hs.26706 ESTs 4.527084 ESTs 4.527084 ESTs 4.527084 ESTs 4.527084 ESTs 4.52708 Bypothetical protein FLJ23560 Hs/064 Hs.28803 Hs.169237 hypothetical protein FLJ23560 fis, clone NT 2.82 3.288 3.21688 Hs/7646 Hs.26106 Hs.26106 erepeat containing 3 final rabe appliens CDNA FLJ12830 fis, clone NT 2.82 3.288 4.527084 Hs.26109 Hs.26708 Hs.	15						
321158	43						
S21170 NS3742				HS.89596			
Sum	:			He 172982			
Substitution				110.172002			
321225 AL080073 Hs.251414 Horno saplens mRNA; cDNA DKFZp564B1462 (f 4.17 4.63 321244 AFG88654 gb;yr76c06.s1 Soares fetal liver spleen 3.80 5.26 321317 Al937050 Hs.6298 KIAA1151 protein 1.00 1.00 1.00 321325 AB033041 Hs.137507 KIAA1215 protein (similar to mouse paladin) 0.44 0.93 321342 AA127984 Hs.222024 transcription factor BMAL2 4.94 4.93 321365 R93443 Hs.271770 ESTs 3.10 4.66 321450 AV05890 Hs.82845 gb;Human 2a12 mRNA for kappa-immunoglobu 2.42 3.35 a21450 W02356 Hs.268980 ESTs 2.26 321498 AV295517 Hs.255436 Hs.268980 ESTs 2.28 3.16 AA703650 Hs.255748 ESTs 2.14 3.94 AV29517 Hs.255436 Hs.268980 Hs.268980 ESTs 2.28 3.1510 AA703650 Hs.255748 ESTs 2.14 3.94 AV29517 Hs.265543 Hs.256514 A019964 Hs.28803 321557 A82964 Hs.268980 Hs.28691 Hs.266514 hs.266514 Hs.266514 hs.26963 Hs.26980 Hs.268980 ESTs 2.28 3.1510 AA703650 Hs.255748 ESTs 2.14 3.94 A689 A21567 AA295430 Hs.25531 Hs.266514 hs.26661 Hs.28803 321567 AA295430 Hs.265614 hs.26661 Hs.28803 321662 AA295430 Hs.266514 hs.266624 Hs.26893 321662 AA295430 Hs.266514 hs.266624 Hs.268980 ESTs 3.06 7.19 321562 AA295430 Hs.266514 hs.266514 hs.26669 Hs.268980 ESTs 3.06 7.19 321567 AA295430 Hs.266514 hs.266514 hs.266514 hs.266514 hs.26669 Hs.268980 ESTs 3.06 7.19 321562 AA295430 Hs.266514 hs.266514 hs.266522 hypothetical protein FLJ11342 4.89 7.82 321628 AW395917 Hs.266514 hs.266514 hs.266021 Hs.26602.r1 Soares retina N2b5HR Homo apil 3.80 7.19 321668 Hs7064 Hs.261601 ESTs Moderately similar to ALU6_HUMAN A 0.47 1.02 321668 Hs7064 Hs.161051 ESTs 2.17 2.45 321688 Hs7064 Hs.167260 ESTs 321602 AA700017 Hs.173737 ras-related C3 botulinum toxin substrate 0.51 1.08 4.57 7.46	50			Hs.226469			7.32
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55 321317 Al937060 Hs.6298 KIÁA1151 protein 1.81 1.55 321318 AB033041 Hs.37597 KIAA1215 protein 1.00 1.00 321325 AB033100 Hs.300646 KIAA protein (similar to mouse paladin) 0.44 0.93 321356 R93443 Hs.271770 ESTs 3.10 4.66 321418 AI739161 Hs.161075 ESTs 2.28 2.54 321420 AJ368667 Hs.132743 ESTs 2.28 2.54 321430 U05890 Hs.132743 ESTs 1.13 0.97 321467 X13075 ESTs 2.26 3.35 321467 X13075 Hs.38540 ESTs 2.46 6.50 321491 H70665 Hs.292549 ESTs 2.46 6.50 321492 W02356 Hs.255748 ESTs 2.28 3.86 321510 AA703650 Hs.255748 ESTs 2.14 3.94 321566 AJ32553							
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321693 AA700017 Hs.173737 ras-related C3 botulinum toxin substrate 0.51 1.08 257 25700 N55160 Hs.167260 ESTs 4.57 7.46				Hs.123158			
321700 N55160 Hs.167260 ESTs 4.57 7.46							
	0.5			Hs.167260		4.57	7.46
	85	321701	AW390923	Hs.42568	ESTs	1.00	1.00

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	321709	N25847	Hs.108923	RAB38, member RAS oncogene family	1.00	1.00
	321710	N35682	Hs.259743	ESTs	2.97	5.26
	321775	AI694875	Hs.202312	Homo sapiens clone N11 NTera2D1 teratoca	1.00	1.00
_	321777	A1637993	Hs.202312	Homo sapiens clone N11 NTera2D1 teratoca	1.68	0.45
5	321779	N42729	Hs.163835	ESTs	0.90	0.90
	321829	D81993	Hs.8966	turnor endothelial marker 8	2.69	3.89
	321846	AA281594	Hs.87902	ESTs	5.11	7.64
	321879	AL109670	Hs.302809	ESTs	6.49	9.58
10	321883	AA426494	Hs.46901	KIAA1462 protein	0.28	0.95
10	321899	N55158	Hs.29468	ESTs	0.39	0.95
	321911	AF026944	Hs.293797	ESTs	6.20	10.76
	321949	R49202	Hs.181694	EST	4.62	10.51
	321955	AI651866	Hs.195689	ESTs	2.89	5.47
	321956	AL110177	Hs.132882	ESTs	0.32	1.25
15	321987	AL133612	Hs.272759	KIAA1457 protein	1.00	1.83
	321991	AL133627	Hs.158923	Homo sapiens mRNA; cDNA DKFZp434K0722 (f	4.00	6.47
	322002	AA328801	Hs.84522	ESTs	2.10	3.48
	322035	AL137517	Hs.306201	hypothetical protein DKFZp564O1278	1.00	1.90
	322044	AW340926		gb:xy51b10.x1 NCI_CGAP_Lu34.1 Homo sapie	3.20	9.67
20	322057	N92197	Hs.154679	synaptotagmin 1	1.55	1.07
	322060	AI341937		gb:qt10e03.x1 NCl_CGAP_GC4 Homo saplens	4.59	7.68
	322070	U80769	Hs.210322	Homo sapiens mRNA for KIAA1766 protein,	2.78	4.52
	322083	AF074982	Hs.226031	ESTs, Highly similar to KIAA0535 protein	3.10	5.52
	322091	AI819863	Hs.106243	ESTs	1.59	1.75
25	322125	R93901		gb:yq16c12.r1 Soares fetal liver spleen	2.06	5.27
	322130	R98978	Hs.117767	ESTs	10.12	16.49
	322147	AF085919	Hs.114176	ESTs	0.94	0.64
	322166	AF085958		gb:yr88b03.r1 Soares fetal liver spleen	4.09	6.67
	322173	H52567		gb:yt85d04.r1 Soares_pineal_gland_N3HPG	3.46	4.85
30	322178	H56535		gb:yt88g03.r1 Soares_pineal_gland_N3HPG	0.44	2.54
	322179	H92891		gb:yt94c02.s1 Soares_pineal_gland_N3HPG	4.52	7.50
	322186	H67346	Hs.269187	ESTs	0.15	0.98
	322196	W87895	Hs.211516	ESTs	2.20	5.04
	322212	AF087995	Hs.134877	ESTs	3.42	4.84
35	322221	AI890619	Hs. 179662	nucleosome assembly protein 1-like 1	0.82	2.14
	322277	Al640193	Hs.226389	ESTs	3.62	3.98
	322278	AF086283		gb:zd46f01.r1 Soares_fetal_heart_NbHH19W	1.00	1.00
	322284	AI792140	Hs.49265	ESTs	0.66	2.76
	322288	AL037273	Hs.7886	pellino (Drosophila) homolog 1	0.71	0.70
40	322320	AF086419		gb:zd78d03.r1 Soares_fetal_heart_NbHH19W	2.02	2.76
	322336	AA308526	Hs.76152	decorin	2.92	4.44
	322339	W17348		gb:zb18c07.x5 Soares_fetal_lung_NbHL19W	8.50	11.56
	322366	AW404274	Hs.122492	hypothetical protein	0.61	1.34
	322372	W25624	Hs.153943	ESTs	7.37	12.07
45	322374	Al394663	Hs.122116	ESTs, Moderately similar to Osf2 [M.musc	4.78	10.50
	322378	AF064819	Hs.201877	DESC1 protein	1.00	1.00
	322388	Al815730	Hs.247474	hypothetical protein FLJ21032	7.09	8.49
	322416	AA223183	Hs.298442	adaptor-related protein complex 3, mu 1	3.20	5.80
	322419	AA248987	Hs.14084	ring finger protein 7	1.64	1.57
50	322425	W37943	Hs.34892	KIAA1323 protein	0.83	. 1.00
	322431	AA069222	Hs.141892	ESTs	3.96	5.22
	322450	AA040131	Hs.25144	ESTs .	5.18	12.67
	322465	AA137152	Hs.286049	phosphoserine aminotransferase	3.41	2.23
	322467	AF116826	Hs.180340	putative protein-tyrosine kinase	1.00	1.30
55	322473	AA744286	Hs.266935	tRNA selenocysteine associated protein	1.75	2.03
	322509	T52172	Hs.302213	ESTs	1.00	2.27
	322523	W80398	Hs.193197	ESTs	2.75	5.49
	322527	AF147359		gb:Homo sapiens full length insert cDNA	1.25	1.27
	322560	AI916847	Hs.270947	ESTs	4.57	8.81
60	322566	W87285	Hs.269587	ESTs	1.00	1.42
	322585	AA837622		gb:zh69c01.r1 Soares_fetal_liver_spleen_	4.18	6.94
	322635	AA679084		gb:zh90h08_r1 Soares_fetal_liver_spleen_	2.40	4.85
	322641	AA007352	Hs.256042	ESTs	2.94	4.64
	322653	AI828854	Hs.258538	striatin, calmodulin-binding protein	0.48	0.38
65	322664	AA011522		gb:zi03g07.r1 Soares_fetal_liver_spleen_	1.92	2.18
	322687	Al110759		gb:AF074666 Human fetal liver cDNA libra	4.14	6.75
	322692	AA018117	Hs.60843	potassium voltage-gated channel, shaker-	3.50	5.00
	322694	Al110872	Hs.279812	PRO0327 protein	1.80	1.72
	322708	AF113674	Hs.283773	clone FLB1727	1.00	3.43
70	322712	AA021328	Hs.23607	hypothetical protein FLJ11109	3.28	3.86
. •	322766	AW068805	Hs.288467	Homo sapiens cDNA FLJ12280 fis, clone MA	1.63	1.53
	322770	AA045796	Hs.122682	ESTs	1.53	1.06
•	322794	Al608591	Hs.38991	S100 calcium-binding protein A2	12.06	1.94
	322810	Al962276	Hs.127444	ESTs	4.09	6.90
75	322818	AW043782	Hs.293616	ESTs	1.20	1.63
	322820	AI377755	Hs.120695	ESTs	0.21	1.93
	322872	AA827228	Hs.126943	ESTs	2.04	1.63
	322882	AW248508	Hs.279727	Homo sapiens cDNA FLJ14035 fis, clone HE	5.26	1.22
	322887	Al986306	Hs.86149	phosphoinositol 3-phosphate-binding prot	2.80	2.24
80	322913	AI733737	Hs.68837	ESTs	2.38	6.61
	322926	AIB25940	Hs.211192	ESTs	4.02	5.79
	322929	Al365585	Hs.146246	ESTs	0.30	1.14
	322968	A1905228	Hs.83484	SRY (sex determining region Y)-box 4	2.06	1.13
	322971	C15953	Hs.212760	hypothetical protein FLJ13649	1.18	2.00
85	322981	AA493252	Hs.159577	ESTs	2.28	2.61
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	322988	C18727	Hs.171941	ESTs	0.39	2.00
	323003	A1733859	Hs.149089	ESTs	3.28	1.00
	323013	AA134042	Hs.191451	ESTs	3.38	5.68
	323025	AL157565	Hs.315369	Homo sapiens cDNA; FLJ23075 fis, clone L	0.06	1.10
5	323032	AW244073	Hs.145946	ESTs	10.18	21.27
	323052	R21124	Hs.85573	Homo sapiens DC29 mRNA, complete cds	1.46	1.90
	323064	AL119341	Hs.49359	Homo sapiens mRNA; cDNA DKFZp547E052 (fr	3.08	5.64
	323098	AI700025	Hs.270471	ESTs	2.31	4.49
	323102	AL119913	Hs.163615	ESTs	5.38	11.64
10	323155	AL135041		gb:DKFZp762K2310_r1 762 (synonym: hmel2)	2.38	5.56
	323176	AW071648	Hs.82101	pleckstrin homology-like domain, family	1.06	1.41
	323191	AA195600	Hs.301570	ESTs	0.73	1.24
	323225	AA205654	Hs.24790	KIAA1573 protein	5.25	11.95
	323232	AA148722	Hs.224680	ESTs	0.45	1.35
15	323266	AW003362	Hs.243886	nuclear autoanligenic sperm protein (his	1.71	1.83
10	323281	A1697556	Hs.292659	ESTs	1.24	3.21
			Hs.86682	Homo sapiens cDNA: FLJ21578 fis, clone C	12.68	15.05
	323283	AA256014	Hs.191501	ESTs	4.42	9.61
	323314	AA226310			2.98	5.93
20	323316	AL134620	Hs.280175	ESTs	1.98	3.30
20	323334	Al336501	Hs.77273	ras homolog gene family, member A		1.00
	323338	R74219	Hs.23348	S-phase kinase-associated protein 2 (p45	1.62 1.00	1.07
	323348	AA233056	Hs.191518	ESTs		1.68
	323351	AA704103	Hs.24049	ESTs	1.43	
25	323359	AA234172	Hs.137418	ESTs	0.34	1.18
25	323360	AA716061	Hs.161719	ESTs	3.01	3.71
	323405	AW139550	Hs.115173	ESTs	1.90	8.81
	323420	A1672386	Hs.263780	ESTs	0.29	1.01
	323434	AW081455	Hs.120219	ESTs	2.27	1.92
••	323445	AA253103	Hs.135569	ESTs, Weakly similar to NEUROD [H.sapien	0.43	0.80
30	323449	AA282865	Hs.284153	Fanconi anemia, complementation group A	3.19	3.85
	323492	H00978	Hs.20887	hypothetical protein FLJ10392	2.70	3.20
	323501	AA182461	Hs.84520	ESTs	2.04	3.31
	323505	Al652287		gb:EST382593 MAGE resequences, MAGK Homo2.	21	3.08
	323515	AA282274	Hs.256083	ESTs	2.69	3.40
35	323541	Al185116	Hs.104613	RP42 homolog	1.20	1.09
	323545	AI814405	Hs.224569	ESTs	1.25	1.55
	323635	R63117	Hs.9691	Homo sapiens cDNA: FLJ23249 fis, clone C	0.27	0.72
	323675	AA984759	Hs.272168	tumor differentially expressed 1	3.70	5.80
	323678	AL042121	Hs.20880	ESTs	3.33	5.10
40	323691	AA317561	Hs.145599	ESTs	1.00	1.00
••	323693	AW297758	Hs.249721	ESTs	2.01	1.54
	323746	AW298611	Hs.12808	MARK	4.11	5.53
	323774	AA329806	Hs.321056	Homo sapiens mRNA; cDNA DKFZp586F1322 (f	2.06	3.70
	323856	AA355264	Hs.267604	hypothetical protein FLJ10450	3.42	8.13
45	323857	T18988	Hs.293668	ESTs	5.97	12.51
-10	323870	AA341774	Hs.129212	ESTs	3.17	4.52
	323876	AL042492	Hs.147313	ESTs	0.36	1.00
	323885	AA344308	Hs.128427	Homo saplens BAC clone RP11-335J18 from	2.31	3.33
	323911	AL043212	Hs.92550	ESTs	4.38	5.41
50	323919	AA862973	Hs.220704		5.80	10.20
50	323972	AIB69964	Hs.182906	ESTs	3.10	5.14
	324005	AA610011	Hs.208021	ESTs	5.34	10.07
	324036	AI472078	Hs.303662	ESTs	1.00	5.03
			Hs.128644	ESTs	0.86	1.00
55	324055	AA528794			0.45	0.91
55	324063	AW292740	Hs.272813	dual oxidase 1 ob:EST94855 Activated T-cells I Homo sap	2.82	5.12
	324072	AA381829	Hs.202473	Homo sapiens cDNA: FLJ22278 fis, clone H	2.40	2.52
	324092	AW269931		Homo sapiens cDNA: FLJ22502 fis, clone H	1.32	4.30
	324095	AW377983	Hs.298140		1.40	1.77
60	324129	Al381918	Hs.285833	Homo sapiens cDNA: FLJ22135 fis, clone H	4.24	6.21
UU	324132	AW504860	Hs.288836	hypothetical protein FLJ12673	6.96	10.69
	324214	AA412395	Hs.225740	ESTs		
	324227	AA295552	Hs.28631	Homo sapiens cDNA: FLJ22141 fis, clone H	0.81	0.53 4.05
	324266	AL047634	Hs.231913	ESTs	2.42	
6 5	324275	AA429088	Hs.98523	ESTs	3.62	5.38
65	324281	AL048026	Hs.124675	ESTs, Weakly similar to T14742 hypotheti	0.14	0.70
	324290	AA432032	Hs.304420	ESTs	3.71	4.34
	324303	AL118754	:	gb:DKFZp761P1910_r1 761 (synonym: hamy2)	0.95	0.91
	324312	Al198841	Hs.128173	ESTs	4.06	5.91
πò	324325	AL138153	Hs.300410	ESTs	5.88	8.25
70	324338	AL138357	Hs.145078	regulator of differentiation (in S. pomb	0.87	1.25
	324341	AW197734	Hs.99807	ESTs, Weakly similar to unnamed protein	1.28	1.00
	324343	AW452016	Hs.293232	ESTs	2.54	3.46
	324371	AA452305	Hs.270319	ESTs	5.85	8.36
	324382	AW502749	Hs.24724	MFH-amplified sequences with leucine-ric	0.76	1.64
75	324384	AA453396	Hs.127656	KIAA1349 protein	2.88	5.69
	324385	F28212	Hs.284247	KIAA1491 protein	1.81	1.99
	324388	Al924963	Hs.306206	hypothetical protein FLJ11215	1.00	1.00
•	324432	AA464510	Hs.152812	ESTs	2.73	217
	324497	AW152624	Hs.136340	ESTs, Weakly similar to unnamed protein	0.71	1.90
80	324510	AI148353	Hs.287425	Homo sapiens cDNA FLJ11569 fis, clone HE	1.00	1.00
- '	324580	AA492588		gb:ng99c08.s1 NCI_CGAP_Thy1 Homo sapiens	2.18	3.50
	324582	AA506935	Hs.132036	ESTs, Weakly similar to ALU1_HUMAN ALU S	5.96	11.36
	324633	AA572994	Hs.325489	ESTs	2.92	4.22
	324640	AW295832	Hs.134798	ESTs, Moderately similar to TTL MOUSE TU	5.48	11.74
85	324675	AW014734	Hs.157969	ESTs	0.39	0.73

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	W	O 02/08	6443			
	324699 324747 324748 324801	AW504732 AA603532 AA657457 AI819924	Hs.21275 Hs.130807 Hs.292385 Hs.14553	hypothetical protein FLJ11011 ESTs ESTs sterol O-acyltransferase (acyl-Coenzyme	0.93 1.57 1.55 1.00	0.93 1.81 1.34 6.56
5	324804 324828 324855 324866	Al692552 AA843926 AW152305 Al541214	Hs.124434 Hs.122364 Hs.46320	gb:wd73f12x1 NCI_CGAP_Lu24 Homo sapiens ESTs ESTs Small proline-rich protein SPRK (human,	1.00 2.00 2.74 1.07	7.53 3.25 3.43 0.95
10	324871 324886 324889 324948 324953	AW297755 AA806794 D31010 AW383618 Al264628	Hs.271923 Hs.131511 Hs.265459 Hs.125428	Homo sapiens cDNA: FLJ22785 fis, clone K ESTs gb:HUML12147 Human fetal lung Homo sapie ESTs, Moderately similar to ALU2_HUMAN A ESTs	1.68 2.56 2.20 5.28 3.37	1.21 5.61 4.65 7.05 5.51
15	324958 324988 325024 325105 325108	AA625076 T06997 F13254 H97109 AA401863	Hs.132892 Hs.121028 Hs.78672 Hs.105421 Hs.22380	protocadherin 20 hypothetical protein FLJ10549 laminin, alpha 4 ESTs ESTs	5.12 2.52 5.24 1.00 1.99	9.81 1.08 10.22 1.00 2.14
20	325114 325146 325149 325187 325228	D83901 Al064690 D61117 Al653682	Hs.315562 Hs.171176 Hs.187646 Hs.197812	ESTS ESTS ESTS ESTS	2.73 1.86 0.42 6.50 6.18	3.17 3.41 0.93 11.31 15.76
25	325235 325328 325340 325367 325373	•			2.64 2.87 0.29 16.56 0.63	4.12 4.42 0.33 24.29 1.22
30	325389 325436 325471 325498				0.88 5.75 8.46 3.32 5.51	1.05 14.14 17.82 6.42 8.28
35	325557 325559 325560 325569 325585				7.48 4.08 4.20 1.10	21.40 6.25 5.24 1.13
40	325587 325597 325639 325685 325686				1.00 2.98 0.78 0.46 0.95	1.00 13.40 0.78 0.66 1.55
45	325735 325739 325740 325792 325819				4.48 0.59 2.42 7.88 4.74	9.20 0.88 6.61 9.83 7.18
50	325883 325895 325925 325932		•		2.02 7.78 2.04 4.18 3.66	2.64 15.98 10.60 7.36 9.03
55	325941 325969 325971 326025 326046				0.61 4.88 0.55 7.21 3.60	0.80 7.42 1.07 14.72 5.98
60	326099 326108 326163 326165 326189		•	•	1.27 3.27 0.45 0.13	1.06 5.70 1.11 0.45
65	326204 326230 326274 326360 326393				5.60 7.00 1.00 9.86 0.52	9.00 12.01 8.09 15.35 0.77
70	326505 326515 326589 326592				1.00 1.24 9.20 2.77	1.42 5.84 13.49 4.01 2.53
75	326605 326692 326693 326720 326742				2.01 1.00 1.00 0.19 2.34	1.00 1.31 0.65 7.20
,	326770 326818 326936 326964	·			0.25 3.09 2.08 0.41	0.83 4.56 3.45 1.70
80	326983 326991 327036 327040 327053				2.02 1.09 1.00 3.05 3.55	3.80 1.20 8.04 4.22 6.31
85	327075				1.59	1.40

	w	O 02/086443		
	327085	0 02,000 0 00	2.50	12.57
	327130		5.38	8.04
	327156		3.74	6.58
5	327220		1.28	1.54
)	327224 327288		6.56 2.61	12.91 5.40
	327321		2.42	3.11
	327332		6.62	10.58
	327361		2.69	4.41
10	327377		2.04	6.72
	327396		2.61	4.50
	327414		1.00 5.91	8.01 9.65
	327442 327467		6.58	18.01
15	327473		3.79	7.48
	327483		4.08	8.87
	327562		0.68	2.86
	327568		1.00	2.00
20	327606		2.06	3.61
20	327611		5.90	14.26
	327642 327654		4.06 1.05	8.74 2.08
	327734		1.00	1.00
	327775		1.46	11.79
25	327796		3.47	5.65
	327840		3.26	6.64
	327940		5.84	15.58
	327984	•	0.36	1.50
30	328004		1.87	1.42
50	328021 328068		0.42 2.83	0.59 4.68
	328100	•	3.04	5.39
	328101		3.54	5.20
0.5	328113		0.72	0.91
35	328157	•	5.58	5.16
	328196		. 5.76	11.13
	328197 328264		5.98 3.11	10.58 4.88
	328299		3.11 2.20	3.06
40 .	328342	·	1.49	1.94
. •	328365		1.00	1.00
	328369		4.40	7.36
	328381		1.86	4.93
45	328451		5.51	7.56
43	328481		0.13 2.71	0.72 3.97
	328500 328530		5.41	7.62
	328600		3.14	10.68
2.2	328608		4.56	8.17
50	328616		2.24	11.91
	328623		3.04	5.46
	328632	•	0.70	1.19
	328664		3.48	6.80
55	328666 328698		10.42 9.68	26.47 14.56
	328700		2.74	10.22
	328708		0.15	0.57
	328735		6.23	`8.91
~	328743		3.62	6.54
60	328806	•	0.22	0.78
	328861		3.68	10.54
	328908 328933		5.42 2.02	16.36 5.29
	328934	•	1.73	4.45
65	328949	•	3.34	5.41
	329005	•	2.88	7.26
	329011	•	2.52 .	3.72
	329033		1.00	1.03
70	329037		5.07	8.16
70	329067	. •	1.98	2.41
	329134		2.24	3.25
	329157 329178		2.30 2.64	11.04 5.02
	329192		6.41	15.27
75	329194		0.31	0.79
_	329204		1.60	3.75
	329224		2.99	6.11
	329228		0.83	0.83
80	329288		0.63	1.01
ou	329337		1.00	1.00
	329541 329560		0.76 1.34	1.68 2.02
	329588		1.68	2.22
0.5	329643		4.18	11.77
85	329703		1.00	1.00

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	**	0 02/00	0770			4
	329764				5.78	15.50
	329816			•	2.09	5.44
				•	3.13	10.77
	329860					
_	329993				7.83	14.21
5	330020				5.58	13.12
-					3.32	5.57
	330036					
	330052				4.31	7.97
	330085				1.34	1.76
					4.70	12.46
	330088					
10	330093				0.44	1.06
10					3.47	4.83
	330100					
	330108				2.14	3.61
	330107				3.17	6.87
	330120				5.61	11.89
15	330123				4.50	12.74
13					1.55	7.62
	330208					
	330263				13.10	23.38
					2.81	4.98
	330300					
	330313			•	3.00	4.41
20	330366				0.67	0.76
20					4.76	11.82
	330372					
	330385	AA449749	Hs.182971	karyopherin alpha 5 (Importin alpha 6)	2.14	2.15
	330397	D14659	Hs.154387	KIAA0103 gene product	0.40	1.15
				raportion delle bronzoi		
	330468	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	1.11	0.94
25	330472	L24203	Hs.82237	ataxia-telangiectasia group D-associated	1.67	1.17
23					0.46	1.07
	330478	L38486	Hs.296049	microfibrillar-associated protein 4		
	330493	M27826	Hs.267319	endogenous retroviral protease	1.07	0.95
				guanine nucleotide binding protein (G pr	0.97	0.96
	330495	M31328	Hs.71642			
	330506	M61906	Hs.6241	phosphoinositide-3-kinase, regulatory su	0.17	3.66
30	330512		Hs.81256	S100 calcium-binding protein A4 (calcium	0.60	1.06
50	•	M80563				
	330537	U19765	Hs.2110	zinc finger protein 9 (a cellular retrov	2.81	2.07
	330547	U32989	Hs.183671	tryptophan 2,3-dioxygenase	3.91	1.49
					1.15	1.03
	330551	U39840	Hs.299867	hepatocyte nuclear factor 3, alpha		
	330568	U56244		(NONE)	2.83	4.79
35		U90437		gb:Human RP1 homolog mRNA, 3'UTR region	2.08	1.54
22	330599					
	330601	U90916	Hs.82845	Homo sapiens cDNA: FLJ21930 fis, clone H	0.89	1.35
	330605	X02419	Hs.77274	plasminogen activator, urokinase	1.87	1.55
					1.83	1.30
	330609	X04741	Hs.76118	ubiquitin carboxyl-terminal esterase L1		
	330617	X53587	Hs.85266	integrin, beta 4	1.54	1.15
40					1.39	1.19
40	330630	X78669	Hs.79088	reticulocalbin 2, EF-hand calcium bindin		
	330644	Y07755	Hs.38991	S100 calcium-binding protein A2	3.83	1.13
	330650	Z68228	Hs.2340	junction plakoglobin	1.25	0.95
				Junctual placogloum		
	330660	AA347868	Hs.139293	ESTs, Weakly similar to ALU7_HUMAN ALU S	15.50	29.07
	330692	AA017045	Hs.6702	ESTs	1.00	1.00
15					0.20	1.35
45	330707	AA133891	Hs.293690	ESTs		
	330715	AA233707	Hs.11571	Homo sapiens cDNA FLJ11570 fis, clone HE	0.12	1.40
				Integrin, beta 8	6.62	5.42
	330717	AA233926	Hs.52620			
	330722	AA243560	Hs.34382	ESTs	1.40	1.65
	330740	AA297746	Hs.22654	Homo sapiens voltage-gated sodium channe	0.27	2.04
50						
50	330742	AA400979	Hs.25691	receptor (calcitonin) activity modifying	0.44	0.90
	330744	AA406142	Hs.12393	dTDP-D-glucose 4,6-dehydratase	0.71	3.23
				Home sealons aDNA Et 113103 for along NT	1.66	1.52
	330751	AA428286	Hs.29643	Homo sapiens cDNA FLJ13103 fis, clone NT		
	330760	AA448663	Hs.30469	ESTs	0.52	0.90
			Hs.274337	hypothetical protein FLJ20666	0.37	0.97
<i></i>	330763	AA450200				
55	330786	D60374	Hs.49136	ESTs, Moderately similar to ALU7_HUMAN A	0.78	0.84
	330790	T48536	Hs.105807	ESTs	0.23	3.17
	330814	AA015730	Hs.265398	ESTs, Weakly similar to transformation-r	0.37	2.07
•	330827	AA040332	Hs.12744	ESTs	1.60	1.00
		AA063037		ESTs	0.93	1.16
~	330844		Hs.66803			
60	330901	AA157818	Hs.267319	endogenous retroviral protease	1.02	1.03
	330931	F01443	Hs.284256	hypothetical protein FLJ14033 similar to	0.24	0.88
					0.08	1.31
	330952	H02855	Hs.29567	ESTs		
	330961	H10998	Hs.7164	a disintegrin and metalloproteinase doma	1.29	1.26
					0.48	0.96
65	330968	H16568	Hs.23748	ESTs MAAAAGE		
65	331014	H98597	Hs.30340	hypothetical protein KIAA1165	0.29	0.74
	331046	N66563	Hs.191358	ESTs	0.99	8.56
				Un annual and a DNA CI 144002 for along UC	1.24	1.00
	331060	N75081	Hs.157148	Homo sapiens cDNA FLJ11883 fis, clone HE		
	331099	R36671	Hs.83937	hypothetical protein	0.75	1.03
		R41408	Hs.21983	ESTs	1.00	2.75
70	331108		NS.21303	E018		
70	331131	R54797		gb:yg87b07.s1 Soares infant brain 1NIB H	6.04	10.68
	331135	R61398	Hs.4197	ESTs	0.80	0.96
	331170.	T23461	Hs.159293	ESTs	2.63	4.29
	331180	T32446	Hs.6640	Human DNA sequence from PAC 75N13 on chr	1.78	2.71
					1.00	3.01
~ ~	331183	T40769	Hs.8469	ESTs		
75	331203	T82310		(NONE)	1.70	3.80
			He garage		1.20	3.19
	331271	AA059347	Hs.82226	glycoprotein (transmembrane) nmb		
	331306	AA252079	Hs.63931	dachshund (Drosophila) homolog	0.31	1.30
		AA281076	Hs.109221	ESTs	2.09	2.41
	331327					
	331341	AA303125	Hs.23240	Homo sapiens cDNA FLJ13496 fis, clone PL	0.72	2.43
80	331359	AA416979	Hs.46901	KIAA1462 protein	0.09	0.91
O.						
	331363	AA421562	Hs.91011	anterior gradient 2 (Xenepus laevis) hom	1.02	0.87
	331378	AA448881	Hs.49282	hypothetical protein FLJ11088	1.03	1.23
	331384	AA456001	Hs.93847	NADPH oxidase 4	1.40	1.00
	331402	AA505135	Hs.44037	ESTs	1.80	3.93
95						
85	331422	F10802	Hs.163628	ESTs, Moderately similar to ALU7_HUMAN	1.65	1.89

	w	O 02/08	6443			
	331490	N32912	Hs.26813	CDA14	2.48	1.73
	331531	N51343	12.20010	gb:yz15g04.s1 Soares_multiple_sclerosis_	0.98	1.68
	331547	N54811		gb:od74f04.s1 NCI_CGAP_Ov2 Homo saplens	3.80	5.75
_	331578	N67960	Hs.249989	ESTs	0.11	0.67
5	331589	N71027	Hs.152618	ESTs	1.09 0.93	1.38 0.76
	331608 331614	N89861 N92293	Hs.112110 Hs.240272	PTD007 protein EST	0.33	1.34
	331668	W69707	Hs.58030	EST	2.24	3.82
	331671	W72033	Hs.194695	ras homolog gene family, member l	1.00	1.24
10	331676	W79834	Hs.58559	ESTs, Weakly similar to rhotekin [M.musc	0.08	1.07
	331681	W85712	Hs.119571	collagen, type III, alpha 1 (Ehlers-Dani	8.72	4.27
	331692	W93592	Hs.152213	wingless-type MMTV integration site fami	0.94 1.57	0.54 1.34
	331717	AA190888 AA191404	Hs.153881 Hs.104072	Homo sapiens NY-REN-62 antigen mRNA, par ESTs	6.80	11.77
15	331718 331811	AA404500	Hs.301570	ESTS	1.10	1.00
10	331820	AA405970	Hs.97996	transcription termination factor, mitoc	0.73	0.59
	331831	AA412031	Hs.97901	EST	2.77	4.08
	331852	AA418988	Hs.98314	Homo sapiens mRNA; cDNA DKFZp586L0120 (f	0.23	0.93
20	331943	AA453418	Hs.21275	hypothetical protein FLJ11011	0.36 1.00	1.88 1.00
20	331969	AA460702	Hs.82772	collagen, type XI, alpha 1 ESTs	3.04	3.87
	331990 332002	AA478102 AA482009	Hs.139631 Hs.105104	ESTS	1.19	0.78
	332027	AA489671	Hs.65641	hypothetical protein FLJ20073	1.27	1.03
	332029	AA489697	Hs.145053	EST8	0.30	1.62
25	332033	AA489840	Hs.251014	EST	2.30	3.70
	332048	AA496019	Hs.201591	ESTs	0.17	0.52
	332071	AA598594	Hs.205293	KIAA1211 protein gb:ae41e11.s1 Gessler Wilms tumor Homo s	1.35 0.19	1.23 2.00
	332074 332083	AA599012 AA600200	Hs.155546	KIAA1080 protein; Golgi-associated, gamm	0.13	1.18
30	332085	AA600353	Hs.173933	nuclear factor I/A	0.30	1.50
-	332125	AA609861	Hs.312447	ESTs	0.22	0.62
	332177	F10812	Hs.101433	ESTs	8.21	18.03
	332180	H03348	Hs.7327	claudin 1	2.27	1.57
35	332185	H10356	Hs.101689	ESTs	0.09 8.05	1.18 5.02
33	332203 332232	H49388 N48891	Hs.317769 Hs.101915	EST Stargardt disease 3 (autosomal dominant)	0.78	0.85
	332240	N54803	Hs.324267	ESTs, Weakly similar to putative p150 [0.96	1.23
	332261	N70294	Hs.269137	ESTs	2.40	3.74
40	332275	R08838	Hs.26530	serum deprivation response (phosphatidyl	0.27	0.75
40	332280	R38100	Hs.146381	RNA binding motif protein, X chromosome	0.39	1.88
	332299	R69250	Hs.21201	nectin 3; DKFZP566B0846 protein	5.24 1.44	12.76 3.18
	332304 332314	R74041 T25862	Hs.101539 Hs.101774	ESTs hypothetical protein FLJ23045	0.68	1.32
	332384	M11433	Hs.101850	retinol-binding protein 1, cellular	1.71	0.88
45	332434	N75542	Hs.289068	Homo sapiens cDNA FLJ11918 fis, clone HE	0.43	0.86
	332445	T63781	Hs.11112	ESTs	0.68	1.00
	332453	L00205	Hs.111758	keratin 6A	31.54	1.00
	332458	M33493	Hs.250700	tryptase beta 1	0.51 0.79	1.00 1.24
50	332504 332525	AA053917 M17252	Hs.15106 Hs.278430	chromosome 14 open reading frame 1 cytochrome P450, subfamily XXIA (steroid	0.73	1.70
50	332530	M31682	Hs.1735	inhibin, beta B (activin AB beta polypep	0.88	0.66
	332535	N20284	Hs.19280	cysteine-rich motor neuron 1	0.22	1.46
	332539	AA412528	Hs.20183	ESTs, Weakly similar to AF164793 1 prote	0.93	1.49
55	332559	M13955	Hs.166189	cytokeratin 2	0.35	1.13
55	332563	N92924	HS.2/440/	protease, serine, 16 (thymus) E1A binding protein p300	1.00 0.36	1.00 1.05
`	332565 332594	AA234896 AA279313	Hs.25272 Hs.3239	methyl CpG binding protein 2 (Rett syndr	0.53	0.59
	332634	S38953	Hs.283750	tenascin XA	0.38	1.16
ė.	332638	AA283034	Hs.50640	JAK binding protein	1.00	1.70
60	332640	AA417152	Hs.5101	protein regulator of cytokinesis 1	6.15	1.16
	332654	AA001296	Hs.288217	hypothetical protein MGC2941	1.50 1.20	2.73 0.91
•	332665 332692	AA223335 AA496035	Hs.63788 Hs.247926	propionyl Coenzyme A carboxylase, beta p gap junction protein, alpha 5, 40kD (con	0.17	1.12
	332716	L00058	Hs.79070	v-myc avian myelocytomatosis viral oncog	1.00	1.44
65	332736	L13773	Hs.114765	myeloid/lymphoid or mixed-lineage leukem	1.00	1.81
7.7	332758	X93921	Hs.296938	dual specificity phosphatase 7	0.53	0.78
	332781	AA233258	Hs.247112	hypothetical protein FLJ10902	1.44	1.56
	332792				1.70 1.85	1.19 2.47
70	332816 332858				1.04	1.57
70	332906				3.48	8.04
	332911				1.00	1.00
	332912				1.06	4.40
75	332922				1.00	1.00
75	332956				0.42 1.96	0.88 6.34
	332959	-: '			0.56	0.99
	332982 332984				0.30	0.78
	332998				1.47	2.01
80	333058				0.47	1.38
	333097				2.14	3.19
	333121				2.76 1.92	3.70 1.21
	333122 333123				1.85	1.39
85	333138				0.47	0.52

	WO 02/086443					PCT/US02/12476
	333139		1.88 0.21	0.84 0.64		
	333140 333221		1.51	1.11		
5	333260 333380		0.75 6.68	1.01 15.75		
•	333387		4.56 5.05	12.61 8.01		
	333512 333524		2.28	3.98	,	•
10	333585 333503		2.31 2.23	1.53 1.17		
10	333604		2.51 0.52	1.58 0.98		
	333618 333627		1.44	1.36		
15	333628 333650		1.90 1.85	1.90 2.10		
10	333678		1.85 2.18	2.35 5.67		•
	333750 333763		1.99	2.60		
20	333767 333768		1.02 1.78	0.96 1.65		
20	333769		2.15 1.46	2.13 2.53		
	333772 333777		1.00	1.42		
25	333846 333884		2.99 0.47	4.50 0.94		
	333887		0.50 0.43	1.00 0.89		
	333891 333892	•	0.51	0.91		
30	333904 333906		0.26 0.55	1.13 0.98		
50	333948	•	1.70 0.37	2.15 1.09		•
	333954 333966	•	8.10	14.30		
35	333968 334061		0.63 4.24	1.38 12.30		
-	· 334094		1.30 4.55	12.03 8.63		
	334113 334161		0.82	1.59		
40	334183 334187		0.47 1.36	0.76 3.70		
	334219		0.69 1.88	1.04 1.70		
	334222 334223	·	4.72	3.14		
45 .	334239 334255		0.7 9 0.45	0.62 1.10		
	334333 334378		1.00 3.98	3.56 5.76		
	334382		1.50	1.31		
50	334492 - 334562		3.59 5.94	4.7 5 15.40		
	334588 334616		8.14 1.55	19.53 1.56		
	334633		5.16	8.07		
55	334648 334787		0.59 3.70	2.13 7.15		
	334866 334891		8.13 0.32	10.60 1.14		
•	334933	•	1.00 4.01	3.84 7.43		
60	334934 334945		1.04	2.96		
	334967 334990		0.29 1.50	1.14 1.39		
	335015		5.88 0.55	18.65 1.75		
65	335093 335120		4.31	8.01		
	335125 335179	•	0.38 1.24	1.97 1.98		
	335188		0.46	1.47 1.42		
70	335211 335288		1.61 0.73 0.20	0.97 0.26		
	335289 335361		0.20 2.18	1.58		
	335379		2.18 0.50 3.64	0.71 14.94		
75	335414 335416		2.93	3.98		
	335496 335497		0.96 1.71	0.91 1.92	Ť	
	335548 335551		1.15 3.22	2.40 10.54		
80	335558		3.42	4.89		
	335586 335619		5.50 2.99	12.75 3.07		
	335620		3.80 0.28	8.29 0.57		
85	335621 335682		0.46	1.17		

	WO 02/086443		
	335686		2.55 3.81
	335755		2.24 1.07
	335784		0.20 0.97
~	335814		1.13 1.48
5	335815		2.45 3.51 1.00 4.16
	335823		0.49 1.70
	335835 335851		1.66 1.39
	335868		2.98 6.43
10	335896		0.98 0.99
	335936		12.10 21.93
	335948		1.00 1.64
	335983		1.00 4.21
4 ~	335995		0.37 1.17
15	336021		1.04 0.84 11.40 23.54
	336034	•	1.19 1.21
	336038		0.54 1.63
	336066 336107		0.95 0.70
20	336205		3.13 6.29
20	336275		3.20 10.10
	336292		2.34 3.09
	336331		1.00 1.00
~~	336419		0.65 0.79 2.33 2.16
25	336632		2.33 2.16 2.55 2.23
	336633		2.55 2.23 2.19 2.03
	336634		2.69 2.48
	336635 336636		2.13 1.83
30	336637		2.43 2.24
50	33663B		2.31 2.03
	336659		0.60 1.31
	336675		0.31 1.18
25	336684		1.50 1.14
35	336694		4.74 7.10 4.43 6.37
	336716		2.20 0.74
	336721 336798		1.64 2.14
	336900	•	6.14 12.73
40	336948		1.00 1.00
. •	337028		1.30 2.09
	337043		4.01 11.53
	337046		1.67 1.84
15	337054		2.78 7.35 7.20 16.14
45	337128		3.45 5.34
	337162 337183	•	5.72 11.41
	337184		3.72 5.90
	337192		1.27 1.06
50	337194		1.88 1.68
	337229		0.22 1.03
	337268		1.00 3.31 3.23 5.14
	337299		3.23 5.14 2.76 3.72
55	337325		5.80 10.42
55	337389		2.06 6.30
	337493 337497		7.88 20.29
	337500		3.80 4.48
	337549		1.66 2.31
60	337603		1.27 8.54
	337605		5.76 7.16
	337671		0.73 0.97 1.54 0.92
	337755		1.54 0.92 5.07 9.73
65	337786 337809		6.18 12.87
05	337862		3.78 12.97
	337871		2.66 8.16
	337958		0.26 1.34
	338008		1.48 1.12
70	338033		2.38 14.59
	338083		0.65 2.16
	338110		1.00 1.61 5.86 8.25
	338112		1.70 1.97
75	338145 338148		8.07 18.19
, 5	338158		1.30 4.55
	338161		2.58 3.57
	338179		1.00 1.00
	338182		3.32 4.63
80	338189		1.00 3.34
	338197		0.99 1.69 4.58 7.62
	338199		6.01 15.85
	338215 338279		0.53 0.95
85	338316		20.58 38.66

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	338322	3.23	7.39
	338357	4.10	11.39
	338359	10.12	21.59
	338366	0.69	1.02
5	338374	0.40	1.18
	338414	0.47	1.06
	338418	6.12	13.86
	338469	3.09	5.11
	338501	6.28	10.32
10	338506	6.97	12.41
	338523	3.10	5.84
	338549	1.70	2.70
	338561	0.79	0.81
	338662	1.72	1.46
15	338671	0.17	0.91
	338676	210	15.86
	338726	1.20	1.09
	338779	0.12	0.57
~~	338804	0.99	1.67
20	338836	1,00	1.00
	338871	4.30	9.81
	338872	5.02	12.81
	338879	0.23	1.12
~-	338937	6.55	12.26
25	338966	1.76	5.42
	338993	1.00	2.40
	339047	5.26	10.81
	339100	5.10	6.88
•	339114	1.00	1.70
30	339121	1.00	3.75
	339170	10.36	19.67
	339229	4.08	13.48
	339264	2.64	3.83
0.5	339293	1.73	1.94
35			•

TABLE 8B shows the accession numbers for those Pkeys in Table 8A lacking unigenelD's. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

PCT/US02/12476

Unique Eos probeset identifier number Pkey: Unique Eos probeset CAT number: Gene cluster number Accession: Genbank accession numbers 45 CAT number Accessions Pkey 187363_1 AW340926 AA249063 N86075 322044 44320_1 Al341937 AW003063 U34725 AA904742 322060 50 X57414 X57415 321430 42705_1 43034_1 321467 X13075 X13076 46779_1 R93901 AF075073 R93902 322125 46861_1 46873_1 H69434 AF085958 H69846 322166 H52567 H52557 AF085970 H52164 322173 55 46882_1 H56535 AF085980 H56712 322178 H92891 AF085982 H92777 46885 1 322179 1615102_1 H84849 H84252 H84260 H86664 H85320 321577 H95531 H95521 H84529 321587 1615333_1 AA070412 AA102346 AA081885 313723 111953_1 60 627492_1 H22544 H46842 Al204929 320997 47271_1 218439_1 322278 W69304 AF086283 W69200 AA625149 AA313030 AA313052 H97463 321687 129439_1 AA665089 AA135130 AA484059 AA102419 AW877765 313883 47422_1 W79150 AF086419 322320 65 322339 814584_1 AI668646 AI734214 W17348 293660_1 AW979268 AA878419 AA431342 AA431628 314648 682222_1 Al308300 Al308296 300201 306897 25196_-2 A1093967 323155 979809_1 AL120701 AL135041 AL121524 70 38927_1 AF147359 T58511 T58560 322527 322585 473768_2 W88919 W89125 1574395_1 Z42308 H23514 300362 82296_1 AA005129 AA679084 AA694399 322635 AA011522 AA702841 AA011691 AA330797 85042_1 322664 75 315454 380580_1 Al239464 Al239473 AA625812 Al208703 322687 37372_1 AF074666 AJ110759 AF090902 AI903735 AA491283 AI694953 AW976903 AA761362 314852 327472_1 307783 697809_1 AI347274 AW844024 AA381722 AA381829 AW963906 AW963902 AA381242 324072 269032_1 AA488472 W27363 AA317053 BE082689 AW967036 BE079872 AW970512 AA280251 Al652287 BE466438 Al650725 AA551854 AA281574 AW571481 80 300527 221345_1 323505 196389_1 315791 403558_1 AA678177 AA677034 AL118754 AA333202 H38001 324303 233842_1 316519 442885_1 AA847835 AA768376 85 300926 333127_1 AA504860 AA504911

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AF068654 AF068656 AF068655

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306015

306063

306065

306104

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330599

331131

331203

331531

331547

332074

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85

H68696

N56929

W81608

AA179868

AA347682

AA411240

AA464716

AA476427

AA513322

AA576453

AA580401

AA897116

AA906316

AA906725

AA910956

AA911861

AA932805

AA936900

AA970223 NOT_FOUND_entrez

15323_-12 U90437 genbank_R54797

NOT_FOUND_entrez

467396_1 AA828597 N54811 genbank_AA599012

genbank_N51343

U56244

R54797

T82310

N51343

AA599012

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TABLE 8C shows the genomic position for those Pkeys in Table 8A lacking unigene ID's and accession numbers. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

5	Pkey: Ref: Strand:	Seque seque	nce of human ct	e 7 digit numbe vromosome 22.		entitled "The DNA
	Nt_position:		tes nucleotide po			
10	Pkey	Ref	Strand	Nt_position		
	332792	Dunham,	l. et.al.	Plus	73381-73768	
1.5	332816	Dunham,	l. et.al.	Plus	359844-360030	
15	332906	Dunham,		Plus	1923101-1923205	
	332911	Dunham,		Plus	1961767-1961858	
	332912 332922	Dunham,		Plus Plus	1962120-1962246 2009620-2009738	
	332956	Dunham, Dunham,		Plus	2510528-2510658	
20	332959	Dunham,		Plus	2518145-2518213	
	333138	Dunham,		Plus	3369205-3369323	
	333139	Dunham,		Plus	3369495-3369571	
	333221	Dunham,		Plus	3978070-3978187	
25	333380 333387	Dunham, Dunham, 1		Plus Plus	4904775-4904846 4910935-4910997	
23	333512	Ounham,		Plus	5560510-5560564	
	333524	Dunham,		Plus	5612620-5612780	
	333585	Dunham,		.Plus	6234778-6234894	
30	333618	Dunham,		Plus	6562391-6562566	
30	333627 333628	Dunham, I Dunham, I		Plus Plus	6620584-6620903 6629004-6629233	
	333650	Dunham, i		Plus	6796852-6797128	
	333678	Dunham,		Plus	7068223-7068288	
25	333750	Dunham, I		Plus	7608165-7608234	
35	333763	Dunham,		Plus	7692491-7692630	
	333767 333768	Dunham, i Dunham, i		Plus Plus	7694407-7694623 7695440-7695697	
	333769	Dunham, I		Plus	7695440-7695697 7696625-7696707	
	333772	Dunham, I		Plus	7706773-7706902	
40 ·	333777	Dunham, I	. et.al.	Plus	7746805-7746916	
	333846	Dunham, I		Plus	8008623-8008757	
	333884 333887	Dunham, I Dunham, I		Plus Plus	8153960-8154161 8154882-8155025	
	333891	Dunham, I		Plus	8156437-8156709	
45	333892	Dunham, I		Plus	8156825-8157001	
	333948	Dunham, I		Plus	8583497-8583627	
	333954 333966	Dunham, I Dunham, I		Plus Plus	6563186-6563335 8655643-8655826	
	333968	Dunham, I		Plus	8681004-8681241	
50	334061	Dunham, I		Plus	9686941-9687077	
	334094	Dunham, I		Plus	9889953-9890105	
	334113	Dunham, I		Plus	10282459-10282597	
	334161 334219	Dunham, I Dunham, I		Plus Plus	10599033-10599180 12716160-12716384	
55	334239	Dunham, I		Plus	13056569-13056693	
	334333	Dunham, 1		Plus	13603544-13603657	
	334378	Dunham, I		Plus	13907239-13907370	
	334382 334562	Dunham, I Dunham, I		Plus Plus	13915866-13916036 14987847-14987940	
60	334588	Dunham, I		Plus	15032740-15032817	
	334616	Dunham, I		Plus	15176123-15176470	
	334633	Dunham, I		Plus	15333206-15333305	
	334866	Dunham, I		Plus	18872214-18872317	
65	334891 334934	Dunham, I Dunham, I		Plus Plus	19299770-19299944 20103970-20104058	
	335015	Dunham, I		_ Plus	20682792-20682945	
	335120	Dunham, I		Plus	21436286-21436384	
	335125	Dunham, I		Plus	21441390-21441471	
70	335179 335188	Dunham, I		Plus	21634405-21634526 21669118-21669328	
70	335211	Dunham, i Dunham, i		Plus Plus	21774611-21774680	
	335361	Dunham, I		Plus	22807292-22807445	
	335379	Dunham, I		Plus	22899306-22899420	
75	335414	Dunham, I		Plus	23235546-23235684	
13	335416 335496	Dunham, I Dunham, I		Plus Plus	23237354-23237465 24164386-24164545	
	335497	Dunham, I		Plus	24167666-24167869	
	335558	Dunham, I	. et.al.	Plus	24740167-24740347	
90	335586	Dunham, I		Plus	24990333-24990497	
80	335686 335784	Dunham, I		Plus	25439839-25439920 25942710-25942792	
	335823	Dunham, I Dunham, I		Plus Plus	26365925-26366004	
	335983	Dunham, I		Plus	27938968-27939070	
05	335995	Dunham, I	. et.al.	Phus	28009044-28009184	
85	336021	Dunham, 1	. et.al.	Plus	28686482-28686559	

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	336034	Dunham, I. et.al.	Ptus	29014404-29014590
	336038	Dunham, I. et.al.	Phys	29022963-29023165
	336107 336632	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	29987731-29987869 983890-985529
5	336633	Dunham, I. et.al.	Plus	985591-986221
	336634 336635	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	986296-986670 987908-988364
	336636 .	Dunham, I. et.al.	Plus	988418-989185
10	336637	Dunham, I. et.al.	Plus	989276-990813
10	336638 336659	Ounham, I. et.al. Dunham, I. et.al.	Plus Plus	991906-993240 1896402-1896478
	336694	Ounham, I. et.al.	Plus	2420546-2420616
	336721	Dunham, I. et.al.	Plus	3371522-3371586
15	336900 336948	Dunham, I. et.al. Dunham, I. et.al.	Pius Pius	10236423-10236523 12692290-12692381
10	337028	Dunham, I. et.al.	Plus	16644817-16644942
	337054	Dunham, I. et.al.	Plus	17821742-17821922 23478943-23479145
	337162 337183	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	23943605-23943696
20	337184	Dunham, I. et.al.	Plus	23973949-23974016
	337268 337299	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	28011979-28012034 29022656-29022775
	337389	Dunham, I. et.al.	Plus	31401509-31401579
25	337493	Dunham, I. et.al.	Plus	33330760-33330981
25	337549 337755	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	34474472-34474531 3971764-3971900
	337809	Dunham, I. et.al.	Plus	4449069-4449193
	337871	Dunham, I. et.al.	Plus Plus	5443027-5443101 6969162-6969270
30	337958 338008	Dunham, I. et.al. Dunham, I. et.al.	. Plus	7697068-7697236
	338033	Dunham, I. et.al.	Plus	8092128-8092271
	338110 338112	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	10384481-10384621 10391398-10391600
	338145	Dunham, I. et.al.	Plus	11386629-11386692
35	338148	Dunham, I. et.al.	Plus	11448985-11449085
	338179 338197	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	12808775-12808833 13638107-13638181
	338279	Dunham, I. et.al.	Plus	16168944-16169091
40	338316 338322	Dunham, I. et.al. Ounham, I. et.al.	Plus Plus	17089711-17089988 17132477-17132547
40	338357	Dunham, I. et.al.	Plus	18062184-18062402
	338359	Dunham, I. et.al.	Plus	18074402-18074501
	338366 338374	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	18252026-18252189 18371200-18371282
45	338414	Dunham, I. et.al.	Plus	19345573-19345660
	338418 338501	Dunham, I. et.al. Ounham, I. et.al.	Plus Plus	19435506-19435596 21244713-21244828
	338506	Dunham, I. et.al.	Plus	21221871-21221953
50	338523	Dunham, I. et.al.	Pius	21509763-21509864 24404720-24404899
30	338662 338804	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	27236005-27236108
	338836	Dunham, I. et.al.	Plus	27792166-27792272
	338879 338937	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	28410653-28410734 29160655-29160725
55	338993	Dunham, I. et.al.	Plus	30077787-30078184
	339047	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	30760793-30760968 31141580-31141765
	339100 339114	Dunham, I. et.al.	Plus	31456454-31456519
60	339121	Dunham, I. et.al.	Plus	31583467-31583536
60	339170 339293	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	32216399-32216527 33223671-33223819
	332858	Dunham, I. et.al.	Minus	1339607-1339397
	332982 332984	Dunham, I. et.al.	Minus Minus	2628296-2628109 2632606-2632457
65	332998	Dunham, I. et.al. Dunham, I. et.al.	Minus .	27.11704-2711565
	333058	Dunham, I. et.al.	Minus	3028925-3028811
	333097 333121	Dunham, I. et.al. Dunham, I. et.al.	Minus	3204124-3204036 3308446-3308358
7 0	333122	Dunham, I. et.al.	Minus	3309596-3309531
7 0 ·	333123	Dunham, I. et.al. Dunham, I. et.al.	Minus Minus	3310817-3310749 3377220-3376309
	333140 333260	Dunham, I. et.al.	Minus	4308400-4308304
	333603	Dunham, I. et.al.	Minus	6466335-6465727
75	333604 333904	Ounham, I. et.al. Dunham, I. et.al.	Minus Minus	6467090-6466768 8217374-8217261
	333906	Dunham, I. et.al.	Minus	8218238-8218063
	334183	Dunham, I. et.al.	Minus	11832582-11832508 11921456-11921205
	334187 334222	Dunham, I. et.al. Dunham, I. et.al.	Minus Minus	12732417-12732289
80	334223	Dunham, I. et.al.	Minus	12734365-12734269
	334255 334492	Dunham, I. et.al. Dunham, I. et.al.	Minus Minus	13200776-13200692 14478333-14478172
	334648	Dunham, I. et.al. Dunham, I. et.al.	Minus	15363301-15363222
85	334787	Dunham, I. et.al.	Minus	16299093-16298937 20078117-20077991
0.0	334933	Dunham, I. et.al.	Minus	20010111-20011331

	11/	O 02/086	112		
	334945	Dunham, I.	et.al.	Minus	20138885-20138637
	334967	Dunham, I.		Minus	20173311-20173218
	334990 335093	Dunham, I. Dunham, I.		Minus Minus	20341159-20341087 21297367-21297214
5	335288	Dunham, I.		Minus	22304275-22303770
	335289	Dunham, I.		Minus	22305950-22305708
	335548 335551	Dunham, I. Dunham, I.		Minus Minus	24662773-24662673 24679828-24678961
4.0	335619	Dunham, I.		Minus	25082677-25082498
10	335620	Dunham, I.		Minus	25092561-25092434
	335621 335682	Dunham, I. Dunham, I.		Minus Minus	25098878-25098767 25421215-25421093
	335755	Dunham, I.		Minus	25763806-25763747
15	335814	Dunham, i.		Minus	26320043-26319845
15	335815 335835	Dunham, I. Dunham, I.		Minus Minus	26320518-26320421 26393311-26393245
	335851	Dunham, I.		Minus	26604863-26604742
	335868	Dunham, I.		Minus	26711437-26711300
20	335896 335936	Dunham, I. Dunham, I.		Minus	26977639-26977558
20	335948	Dunham, I.		Minus Minus	27360474-27360400 27555924-27555788
	336066	Dunham, I.		Minus	29241080-29240842
	336205	Dunham, I.		Minus	30477456-30477311
25	336275 336292	Dunham, I. Dunham, I.		Minus Minus	32086675-32086536 32818035-32817927
	336331	Dunham, I.		Minus	33594527-33594371
	336419	Dunham, I.		Minus	34052568-34052445
	336675 336684	Dunham, I. Dunham, I.		Minus Minus	2020758-2020664 2158060-2157993
30	336716	Dunham, I.		Minus	3259952-3259862
	336798	Dunham, I.		Minus	5888954-5888757
	337043 337046	Dunham, I. Dunham, I.		Minus Minus	17407330-17407251 17610892-17610821
	337128	Dunham, I.		Minus	22215251-22215034
35	337192	Dunham, I.		Minus	24591853-24591771
	337194 337229	Dunham, I. o Dunham, I. o		Minus Minus	24610510-24610359 26716579-26716481
	337325	Dunham, I.		Minus	30015948-30015800
40	337497	Dunham, I.		Minus	33371317-33371258
+0	337500 337603	Dunham, I. o Dunham, I. o		Minus Minus	33376212-33376158 1299296-1299194
	337605	Dunham, I.		Minus	1346555-1346397
	337671	Dunham, i.		Minus	3260634-3260547
45	337786 337862	Dunham, I. o Dunham, I. o		Minus Minus	4133203-4133081 5347658-5347550
	338083	Dunham, I.		Minus	9318438-9318301
•	338158	Dunham, I. e		Minus	11794465-11794343
	338161 338182	Dunham, I. e Dunham, I. e		Minus Minus	12124716-12124658 12824919-12824827
50	338189	Dunham, I.		Minus	12878594-12878478
	338199	Dunham, I. e		Minus	13760865-13760780
	338215 338469	Dunham, I. e Dunham, I. e		Minus Minus	14055447-14055355 20520387-20520242
	338549	Dunham, I. e		Minus	22049171-22049081
55	338561	Dunham, I. e		Minus	22311966-22311856
•	338671 338676	Dunham, I. e Dunham, I. e		Minus Minus	24508421-24508346 24637427-24637369
	338726	Dunham, I. e		Minus	25926206-25925618
50	338779	Dunham, I. e		Minus	27030151-27029795
JU	338871 338872	Dunham, I. e Dunham, I. e		Minus Minus	28301708-28301611 28300921-28300790
	338966	Dunham, I. e		Minus	29614876-29614749
	339229	Dunham, I. e		Minus	32722330-32722199
55	339264 325228	Dunham, I. e 6381940	t.a!. Plus	Minus 2630-2694	32975145-32975053
,,,	325235	6381943	Minus	162154-162	264
	329588	3962484	Ptus	1169-1619	
	329560 329541	3962491 3983503	Ptus Minus	2095-2990 2765-3059	
70	325328	5866875	Plus	86780-86854	1
	325340	6017033	Minus	166656-1668	319
	325373	5866920	Minus	1136686-113	
	325367 325389	5866920 5866921	Minus Plus	922881-9229 239672-2397	
75	325436	5866939	Minus	29778-29907	
	325498	5866967	Plus	173372-1739	
	325471 325557	6017034 6056302	Minus Plus	289268-2893 50921-51050	
	325559	6249595	Minus	118590-119	
30	325560	6249595	Minus	133794-1339	981
	325569 325587	6249599 6682462	Plus Plus	79927-80217 126724-1269	
	325585	6682462	Plus Plus	73476-73574	
35	325597	5866992	Plus	1065020-106	5089
20	325639	5867002	Plus	253525-2536	808

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	325739 325740	5867038 5867038	Minus Minus	205138-205269 207533-207690				
	325792	6469828	Minus	1018-1176				
5	325735	6552447 6682468	Minus Plus	269122-269190 117397-117483				
,	325685 325686	6682468	Plus	118337-118439				
	325819	6682490	Minus	130314-130370				
	329764 329703	6048195 6065793	Minus Minus	109733-109958 139994-140138				
10	329643	6448539	Plus	53403-53537				
	329816 329860	6624888 6687260	Minus Minus	70296-70423 163474-163605				
	325883	5867087	Pius	22498-22663				
15	325895	5867097	Plus	358317-358476				
13	325925 325932	5867124 5867127	Plus Plus	115749-115962 7369-7441				
	325941	5867133	Minus	64228-64402				
	325969 325971	5867153 5867153	Plus Plus	101911-102081 105841-106035				
20	329993	4567166	Minus	101307-101434				
	330020	6671887	Plus	172397-172491				
	326163 326274	5867168 5867171	Minus Minus	7831-8035 410289-410404				
25	326025	5867176	Plus	70854-70915				
25	326046 326099	5867182 5867186	Minus Minus	62668-62825 661381-661510				
	326108	5867187	Minus	23784-23903				
	326165	5867208	Minus	62787-62929				
30	326189 326204	5867212 5867218	Pius Minus	69288-69413 148088-148200				
	326230	5867230	Minus	301868-301972				
	330052 330036	4567182 6042048	Plus Plus	352560-352963 117120-117216				
	326360	5867293	Plus	13627-13844				
35	326589	5867320	Plus	22760-22919				
	326393 326505	5867341 5867435	Plus Minus	41702-41841 8818-8949				
	326515	5867439	Plus	36683-36809				
40	326592 330107	6138928 6015249	Plus Minus	23689-23828 100091-100282				
••	330106	6015249	Minus	99443-99778				
	330100 330093	6015253 6015278	Plus Plus	21166-21301 1043-1199				
,	330088	6015293	Plus	37517-37638				
45 -	330085	6015302	Minus	59613-59770				
	330120 330123	6671864 6671869	Minus Minus	127553-127656 35311-35406				
	326742	5867611	Minus	95187-95248				
50	326605 326818	5867637 6117831	Plus Minus	24656-24749 15199-15309				
J 0	326720	6552456	Plus	84525-84677				
	326770 326692	6598307 6682502	Minus Plus	513603-513668 117697-117899				
	326693	6682502	Minus	335002-335095				
55	326983	5867657	Minus	16023-16581				
	326991 326936	5867660 6004446	Plus Minus	18147-18339 10217-10357				
	326964	6469836	Plus	75340-75456				
60	327040 327053	6531965 6531965	Plus Plus	783670-783817 2247267-2247437				
•	327075	6531965	Plus	4041318-4041431				
	327085 327036	6531965 6531965	Plus	4734947-4735069 319951-320040				
	327030	6531976	Plus Plus	20247-22343				
65	327156	5866841	Minus	2462-2620				
	327288 327332	5867481 · 5867516	Plus Minus	48583-48773 56361-56532				
	327220	5867525	Minus	65701-65781				
70	327224 327321	5867534 6249562	Plus Minus	188468-188544 99745-99836				
, 0	327361	6552412	Minus	61013-62130				
	327396	5867743	Plus	8702-8820				
	327414 327442	5867750 5867759	Plus Plus	102461-102586 111483-111618				
75	327467	5867772	Plus	88030-88151				
	327473 327483	5867775 5867783	Plus Plus	75101-75181 181573-181662				
	327377	5867793	Minus	37610-37676				
80	327562	5867804	Minus	343989-344474				
ou	327568 327608	5867811 6004463	Minus Plus	46152-46287 200262-200495				
	327611	5867868	Minus	175063-175392				
_	327642 327654	5867891 5867910	Minus Minus	2513-2743 97564-97710				
85	327734	5867940	Minus	31003-31583				

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	327775	5867964	Minus	130791-130871
	327796	5867982	Plus	85267-85405
	327840	6249578	Minus	73065-73206
_	330208	6013599	Pius	66517-66931
5	330263	6671884	Minus	101503-101634
	328004	5867993	Minus	157407-157887
	328101	5868020	Plus	289920-290014
	328100	5868020	Minus	263545-263635
10	328113	5868024	Minus	80378-80491
10	328157	5868064	Plus	73326-73615
	328196	5868080	Minus	16551-16729
	328197	5868081	. Minus	42133-42438
	327940	5868197	Minus	95240-95428
15	327984	5868216	Plus	66611-66677
15	328021	5902482	Plus	713478-714590
	328068	6117819	Plus	253903-254022
	328264	6381912	Plus	55086-55404
	330300	2905862	Minus	3246-3302
20	328608	5868222	Minus	87770-87953 38889-40010
20	328600	5868229	Minus	293920-294224
	328616 328623	5868239 5868246	Plus Minus	120020-120126
	328623 328632	5868247	Plus	76734-76853
	328666	5868254	Minus	778-901
25	328698	5868264	Minus	625555-625633
23	328700	5868264	Plus	764089-764203
	328708	5868271	Minus	68114-68854
•	328735	5868289	Plus	89389-89455
	328743	5868289	Plus	274638-274726
30	328806	5868324	Plus	29408-29684
	328299	5868366	Minus	149708-149889
	328342	5868383	Plus	59955-60094
	328365	5868387	Minus	270724-270798
	328369	5868388	Plus	75371-75583
35	328381	5868392	Plus	662758-662848
	328451	5868425	Minus	217275-217336
	328481	5868449	Minus	8987-9180
	328500	5868464	Plus	59098-59481
40	328530	5868482	Plus	334973-335406
40	328664	6004473	Plus	1193739-1193866
	328861	6381928	Minus	108317-108403
	328908	5868493	Plus	117002-117059
	328933	5868500	Plus	771755-771889
45	328934	5868500 ·	Plus	846342-846448
43	328949	6456765	Minus	43552-43619
	330313	6042030	Minus	33642-33775
	329005	5868542	Plus	85470-85673 151837-151914
	330366 330372	2944106 6580495	Plus Minus	317461-317688
50	329033	5868561	Minus	5390-5479
J U	329033	5868562	Minus	32466-32562
	329067	5868591	Minus	146417-147652
	329134	5868679	Plus	29959-30018
	329157	5868687	Minus	145940-146155
55	329178	5868704	Plus	179177-179463
	329192	5868716	Plus	166936-167020
	329194	5868716	Minus	304450-304559
	329204	5868720	Minus	3050-3190
	329224	5868728	Plus	27422-27664
60 -	329228	5868728	Minus	50118-50287
	329288	5868771	Plus	25554-26299
	329337	5868806	Minus	467155-467222
	329011	6682532	Plus	48658-48741

TABLE 9A: Potential Therapeutic, Diagnostic and Prognostic targets for Therapy of Lung Cancer

Table 9A shows about 1312 genes up-regulated in lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granufomatous and carcinoid tumors) relative to normal body tissues. These genes were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array.

Table 9B show the accession numbers for those Pkey's lacking UnigenelD's for table 9A. For each probaset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarly using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 9C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 9A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

15 Pkey:

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number ExAccn:

Unigene number UnigenelD: Unigene Title: Unigene gene title

5

10

20

Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the R1:

average of normal lung samples

(including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples

	R2:	Averag	e of non-malig	mant lung disease samples (including bronchitis, emphysema	ı, fibrosis,	atelectasis, asth
	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
	400195		-	NM_007057*:Homo sapiens ZW10 interactor	1.00	1.00
25	400205			NM_006265*:Homo sapiens RAD21 (S. pombe)	15.80	396.00
	400220			Eos Control	2.28	2.84
	400277			Eos Control	7.68	9.72
	400285			Eos Control	1.00	1.00
20	400288	X06256	Hs.149609	integrin, alpha 5 (fibronectin receptor,	1.04	2.24
30	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	132.45	4.00
	400298	AA032279	Hs.61635	six transmembrane epithelial antigen of	43.86	74.00 1.00
	400301	X03635	Hs.1657	estrogen receptor 1	1.00 1.75	1.65
	400303	AA242758	Hs.79136	LIV-1 protein, estrogen regulated	0.87	1.80
35	400328	X87344	Hs.180062	transporter 2, ATP-binding cassette, sub	156.55	253.00
33	400419 400512	AF084545		Target NM_030878*:Homo sapiens cytochrome P450,	1.00	2.00
	400512	AF242388		lengsin	3.67	87.00
	400560	AF242300		NM_030878*:Homo sapiens cytochrome P450,	1.00	1.00
	400564			NM_002425:Homo sapiens matrix metallopro	20.26	45.00
40	400665			NM_002425:Homo sapiens matrix metallopro	1.36	1.07
.0	400666			NM_002425:Homo sapiens matrix metallopro	3.26	3.22
	400749			NM_003105*:Homo sapiens sortilin-related	1.00	91.00
	400763			Target Exon	7.63	24.00
	401027			Target Exon	1.00	1.00
45	401093			C12000586*:glj6330167[dbj]BAA86477.1] (A	1.00	155.00
	401203			Target Exon	1.00	86.00
	401212			C12000457*:gi 7512178 pir [T30337 polypr	1.00	400.00
	401411			ENSP00000247172*:HYPOTHETICAL 126.2 kDa	1.00	72.00
~^	401435			C14000397*:gij7499898[pirl]T33295 hypoth	1.00	64.00
50	401464	AF039241		histone deacetylase 5	3.82	49.00
	401714			ENSP00000241802*:CDNA FLJ11007 FIS, CLON	2.02	40.00
	401747			Homo sapiens keratin 17 (KRT17)	128.43	68.00
	401760			Target Exon	1.74	35.00
<i>5 </i>	401780			NM_005557*:Homo sapiens keratin 16 (foca	26.47	10.50
55	401781			Target Exon	10.33	4.61
	401785			NM_002275*:Homo sapiens keratin 15 (KRT1	4.13 1.44	2.70 2.10
	401797			Target Exon	1.41	1.86
	401961 401985	AF053004		NM_021626:Homo saplens serine carboxypep class I cytokine receptor	1.00	177.00
60	401994	AITU33004		Target Exon	61.84	47.00
00	402075			ENSP00000251056*:Plasma membrane calcium	1.00	1.00
	402260			NM_001436*:Homo sapiens fibriliarin (FBL	1.58	1.39
	402265			Target Exon	2.09	35.00
	402297	-		Target Exon	1.00	92.00
65	402408			NM_030920*:Homo sapiens hypothetical pro	28.87	13.00
	402420			C1000823*:gi]10432400 emb[CAC10290.1] (A	1.00	1.44
	402674			Target Exon	7.44	243.00
	402802			NM_001397:Homo sapiens endothelin conver	1.00	70.00
- 0	402994			NM_002463*:Homo sapiens myxovirus (influ	1.37	1.43
70	403137			NM_005381*:Homo sapiens nucleolin (NCL),	1.00	19.00
	403306	NM_006825		transmembrane protein (63kD), endoplasmi	1.00	43.00
	403329			Target Exon	1.00	61.00
	403381			ENSP00000231844*:Ecotropic virus integra	1.00	119.00
75	403478			NM_022342:Homo sapiens kinesin protein 9	28.13	136.00
13	403485			C3001813*:gi 12737279 ref XP_012163.1 k	20.23 6.30	76.00 29.33
	403627			Target Exon	1.30	35.00
	403715 404044			Target Exon ENSP00000237855*:DJ398G3.2 (NOVEL PROTE)	1.00	54. 00
	404044			NM_016020*:Homo sapiens CGI-75 protein (14.29	91.00
80	404076			C8000950:gi 423560 pir A47318 RNA-bindi	1.00	1.00
50	404140			NM_006510:Homo saplens ret finger protei	1.42	1.44
	404140			ENSP00000244562:NRH dehydrogenase (quino	1.00	54.00
	404185			Target Exon	1.00	117.00
	404210			NM_005936:Homo saplens myeloid/lymphoid	5.93	13.77
85	404253			NM_021058*:Homo sapiens H2B histone fami	1.00	1.00

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	404287		•	C6001909:gij704441 dbjjBAA18909.1] (D298	29.71	42.00
	404298			C6001238*:gij121715 sp P26697 GTA3_CHICK	1.30	1.00
	404347 404440			Target Exon NM_021048:Homo sapiens melanoma entigen,	1.00 1.00	1.00 15.00
5	404721			NM_005596*:Homo sapiens nuclear factor I	1.00	60.00
•	404794	NM_000078		cholesteryl ester transfer protein, plas	1.07	1.38
	404854			Target Exon	1.61 1.00	2.01 1.00
	404877 404927			NM_005365:Homo sapiens melanoma antigen, Target Exon	1.00	1.00
10	404996			Target Exon	1.00	1.00
	405449			CY000047*:gi[11427234 ref]XP_009399.1] z	1.00	1.00
	405568 405572			NM_031413*:Homo sapiens cat eye syndrome Target Exon	1.00 0.76	78.00 1.14
	405646			C12000200:gi 4557225 ref NP_000005.1 al	1.01	1.28
15	405676	BE336714		cytochrome c-1	1.13	2.89
	405770			NM_002362:Homo saplens melanoma antigen,	45.52 1.99	37.00 1.99
	405932 406137			C15000305:gij3806122 gbjAAC69198.1 (AF0 NM_000179*:Homo sapiens mutS (E. coli) h	2.77	2.38
	406360			Target Exon	1.00	35.00
20	406399			NM_003122*:Homo sapiens serine protease	1.00	39.00
	406467 406621	X57809	Hs.181125	Target Exon immunoglobulin lambda locus	1.00 1.41	1.00 1.74
	406642	AJ245210	16.101125	gb:Homo sapiens mRNA for immunoglobulin	2.16	3.91
05	406663	U24683	Hs.293441	immunoglobulin heavy constant mu	2.07	2.93
25	406671	AA129547	Hs.285754	met proto-oncogene (hepatocyte growth fa	15.00 0.98	51.00 3.09
	406673 406676	M34996 X58399	Hs.198253 Hs.81221	major histocompatibility complex, class Human L2-9 transcript of unrearranged im,	1.30	1.53
	406678	U77534		gb:Human clone 1A11 immunoglobulin varia	1.33	1.45
20	406685	M18728	11- 070000	gb:Human nonspecific crossreacting antig	1.46 8.61	2.85 8.50
30	406687 406690	M31126 M29540	Hs.272822 Hs.220529	pregnancy specific beta-1-glycoprotein 9 carcinoembryonic antigen-related cell ad	226,37	350.00
	406698	X03068	Hs.73931	major histocompatibility complex, class	1.01	2.52
	406815	AA833930	Hs.288036	tRNA isopentenylpyrophosphate transferas	20.25 0.75	32.00 1.91
35	406851 406964	AA609784 M21305		major histocompatibility complex, class gb:Human alpha satellite and satellite 3	38.15	1114.00
55	406967	M24349		gb:Human parathyroid hormone-like protei	1.00	1.00
	406974	M57293		gb:Human parathyroid hormone-related pep	1.00	1.00
	407103 407128	AA424881 R83312	Hs.256301 Hs.237260	hypothetical protein MGC13170 EST	1.77 1.00	1.10 1.00
40	407137	T97307	110,201200	gb:ye53h05.s1 Soares fetal liver spleen	142,70	135.00
	407168	R45175	Hs.117183	ESTs .	2.16	18.00
	407239 407242	AA076350 M18728	Hs.67846	leukocyte immunoglobulin-like receptor, gb:Human nonspecific crossreacting antig	1.10 1.12	1.57 2.85
	407244	M10014	Hs.75431	fibrinogen, gamma polypeptide	3.24	15.38
45	407289	AA135159	Hs.203349	Homo sapiens cDNA FLJ12149 fis, clone MA	3.53	3.68
	407300 407366	AA102616 AF026942	Hs.120769 Hs.271530	gb:zn43e07.s1 Stratagene HeLa cell s3 93 gb:Homo sapiens cig33 mRNA, partial sequ	19.74 0.06	73.00 8.25
	407378	AA299264	Hs.57776	ESTs, Moderately similar to I38022 hypot	1.00	26.00
50	407430	AF169351		gb:Homo sapiens protein tyrosine phospha	1.00 1.00	25.00 75.00
50	407453 407577	AJ132087 AW131324	Hs.246759	gb:Homo sapiens mRNA for axonemal dynein hypothetical protein MGC12538	1.00	1.00
	407634	AW016569	Hs.136414	UDP-GlcNAc;betaGal beta-1,3-N-acetylgluc	111.20	228.00
	407710	AW022727	Hs.23616	ESTs	1.00 1.89	28.00 1.31
55	407720 407746	AB037776 AK001962	Hs.38002	KIAA1355 protein hypothetical protein FLJ11100	1.00	1.00
	407756	AA116021	Hs.38260	ubiquitin specific protease 18	4.51	5.00
	407758	D50915	Hs.38365	KIAA0125 gene product ESTs, Moderately similar to PURKINJE CEL	1.00 0.97	28.00 1.14
	407782 407788	AA608956 BE514982	Hs.112619 Hs.38991	S100 calcium-binding protein A2	7.88	3.83
60	407790	AI027274	Hs.288941	Homo sapiens cDNA FLJ14866 fis, clone PL	3.63	42.00
	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon ESTs	89.96 173.91	109.00 108.00
	407839 407944	AA045144 R34008	Hs.161566 Hs.239727	desmocollin 2	111.30	70.00
~=	408000	L11690	Hs.620	bullous pemphigoid antigen 1 (230/240kD)	151.17	8.00
65	408031 408063	AA081395 BE086548	Hs.42173 Hs.42346	Homo sapiens cDNA FLJ10366 fis, clone NT calcineurin-binding protein calsarcin-1	9.91 195.78	93.00 231.00
	408070	AW148852	113.42040	gb:xf05d05.x1 NCI_CGAP_Bm35 Homo sapien	1.00	1.00
	408101	AW968504	Hs.123073	CDC2-related protein kinase 7	37.84	61.00
70	408122	A1432652 AA297567	Hs.42824 Hs.43728	hypothetical protein FLJ10718 hypothetical protein	0.85 5.88	1.71 7.91
70	408212 408243	Y00787	Hs.624	interleukin 8	4.27	9.98
	408349	BE546947	Hs.44276	homeo box C10	3.79	3.46
	408353 408354	BE439838 Al382803	Hs.44298 Hs.159235	mitochondrial ribosomal protein S17 ESTs	1.88 1.00	1.65 73.00
75	408369	R38438	Hs.182575	solute carrier family 15 (H??? transport	1.41	16.50
	408380	AF123050	Hs.44532	diubiquitin	15.19	37.22
	408482 408522	NM_000676 Al541214	Hs.45743 Hs.46320	adenosine A2b receptor Small proline-rich protein SPRK [human,	1.65 1.98	1.19 1.24
	408536	AW381532	Hs.135188	ESTs	1.55	1.50
80	408545	AW235405	Hs.253690	ESTs	1.00	1.00
	408572 408633	AA055611 AW963372	Hs.226568 Hs.46677	ESTs, Moderately similar to ALU4_HUMAN A PRO2000 protein	1.00 107.16	44.00 56.00
	408660	AA525775	. 1007.011	ESTs, Moderately similar to PC4259 femi	1.00	1.00
85	408761	AA057264	Hs.238936	ESTs, Weakly similar to (defline not ava	52.24 3.05	141.00 109.00
UJ	408771	AW732573	Hs.47584	potassium voltage-gated channel, delayed	3.03	103.00

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	408783	AF192522	Hs.47701	NPC1 (Niemann-Pick disease, type C1, gen	1.02	1.07
	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,	41.19	61.00
	408805	H69912	Hs.48269	vaccinia related kinase 1	24.67	45.00
5	408841	AW438865	Hs.256862	ESTs	1.00	58.00 89.00
3	408873 408908	AL046017 BE296227	Hs.182278 Hs.250822	calmodulin 2 (phosphorylase kinase, delt serine/threonine kinase 15	1.00 7.76	1.00
	408992	AA059325	Hs.71642	guanine nucleotide binding protein (G pr	1.00	1.00
	408996	Al979168	Hs.344096	glycoprotein (transmembrane) nmb	3.71	5.50
4.0	409015	BE389387	Hs.49767	NM_004553:Homo sapiens NADH dehydrogenas	1.44	1.24
10	409038	T97490	Hs.50002	small inducible cytokine subfamily A (Cy	4.28	5.32
	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119	112.42 1.00	195.00 17.00
	409077 409093	AA401369 BE243834	Hs.190721 Hs.50441	ESTs CGI-04 protein	2.02	1.93
	409103	AF251237	Hs.112208	XAGE-1 protein	80.44	40.00
15	409142	AL136877	Hs.50758	SMC4 (structural maintenance of chromoso	14.87	6.00
	409187	AF154830	Hs.50966	carbamoyl-phosphate synthetase 1, mitoch	1.00	1.00
	409228	AI654298	Hs.271695	ESTs, Weakly similar to 2109260A B cell	1.22	1.00
	409234	AI879419	Hs.27206	ESTs	1.00	1.00
20	409268	AA625304	Hs.187579 Hs.22972	ESTs	11.90 1.00	23.00 1.00
20	409269 409361	AA576953 NM_005982	Hs.54416	hypothetical protein FLJ 13352 sine oculis homeobox (Drosophila) homolo	168.91	35.00
•	409404	BE220053	Hs.129056	ESTs	1.00	1.00
	409420	Z15008	Hs.54451	laminin, gamma 2 (nicein (100kD), kalini	79.74	96.00
05	409430	R21945	Hs.346735	splicing factor, arginine/serine-rich 5	1.45	2.10
25	409446	Al561173	Hs.67688	ESTs	1.00	4.00
	409506	NM_006153	Hs.54589	NCK adaptor protein 1	3.97 15.98	28.00 141.00
	409522 409582	AA075382 AA401369	Hs.190721	gb:zm87b03.s1 Stratagene ovarian cancer ESTs	1.00	17.00
	409632	W74001	Hs.55279	serine (or cysteine) proteinase inhibito	292.12	79.00
30	409705	M37762	Hs.56023	brain-derived neurotrophic factor	1.00	82.00
	409719	A1769160	Hs.108681	Homo sapiens brain turnor associated prot	1.00	1.00
	409731	AA125985	Hs.56145	thymosin, beta, identified in neuroblast	0.12	18.12
	409744	AW675258	Hs.56265	Homo sapiens mRNA; cDNA DKFZp586P2321 (f	20.75 22.46	51.00 15.80
35	409757 409866	NM_001898 AW502152	Hs.123114	cystatin SN gb:UI-HF-BR0p-air-f-11-0-UI.r1 NIH_MGC_5	1.00	1.00
55	409893	AW247090	Hs.57101	minichromosome maintenance deficient (S.	1.50	1.09
	409902	Al337658	Hs.156351	ESTs	25.92	50.00
	409935	AW511413	Hs.278025	ESTs	2.63	2.11
40	409956	AW103364	Hs.727	inhibln, beta A (activin A, activin AB a	2.17	4.01
40	409958	NM_001523	Hs.57697	hyaluronan synthase 1	0.91	2.07
	410001 410032	AB041036	Hs.57771	kæliikrein 11 gb:RC3-BT0319-120200-014-a09 BT0319 Homo	1.04 1.00	2.28 58.00
	410032	BE065985 AB020725	Hs.58009	KIAA0918 protein	1.00	34.00
	410044	BE566742	Hs.58169	highly expressed in cancer, rich in teuc	1.00	1.00
45	410048	W76467	Hs.58218	proline oxidase homolog	1.03	1.44
	410076	T05387	Hs.7991	ESTs	1.12	1.50
	410102	AW248508	Hs.279727	Homo sapiens cDNA FLJ14035 fis, clone HE	9.89	1.00
	410153	BE311926	Hs.15830	hypothetical protein FLJ 12691 hypothetical protein FLJ 10514	1.00 1.00	1.00 1.00
50	410166 410193	AK001376 AJ132592	Hs.59346 Hs.59757	zinc finger protein 281	42.01	51.00
20	410274	AA381807	Hs.61762	hypoxia-inducible protein 2	1.72	1.32
	410309	BE043077	Hs.278153	ESTs	1.00	2.00
	410340	AW182833	Hs.112188	hypothetical protein FLJ 13149	32,08	75.00
55	410348	AW182663	Hs.95469	ESTs	1.00	1.00
99	410407	D31382	Hs.63287 Hs.63325	carbonic anhydrase IX transmembrane protease, serine 4	1.40 4.30	1.11 2.03
	410438	AB037756	Hs.45207	hypothetical protein KIAA1335	1.00	18.00
	410553	AW016824	Hs.255527	hypothetical protein MGC14128	1.34	1.04
	410555	W27235	Hs.64311	a disintegrin and metalloproteinase doma	23.99	1.41
60	410561	BE540255	Hs.6994	Homo sapiens cDNA: FLJ22044 fis, clone H	10.04	1.00
	410681	AW246890	Hs.65425	calbindin 1, (28kD)	10.88	18.92
	410781 411027	A1375672 AF072099	Hs.165028 Hs.67846	ESTs leukocyte immunoglobulin-like receptor,	1.00 1.62	57.00 3.78
	411074	X60435	Hs.68137	adenylate cyclase activating polypeptide	1.00	1.15
65	411089	AA456454	110.00	cell division cycle 2-like 1 (PITSLRE pr	1.56	1.58
	411152	BE069199		gb:QV3-BT0379-010300-105-g03 BT0379 Homo	1.00	84.00
	411248	AA551538	Hs.334605	Homo sapiens cDNA FLJ14408 fis, clone HE	1.82	1.45
	411252	AB018549	Hs.69328	MD-2 protein	7.32	12.74
70	411263	BE297802	Hs.69360 Hs.289082	kinesin-like 6 (mitotic centromere-assoc GM2 ganglioside activator protein	3.44 1.35	2.55 2.02
70	411365 411402	M76477 BE297855	Hs.69855	NRAS-related gene	1.00	46.00
	411573	AB029000	Hs.70823	KIAA1077 protein	11.40	11.35
	411579	AC005258	Hs.70830	U6 snRNA-associated Sm-like protein LSm7	1.08	1.90
75	411617	AA247994	Hs.90063	neurocalcin delta	1.74	2.57
75	411732	AA059325	Hs.71642	guarrine nucleotide binding protein (G pr	1.02	1.00
	411773	NM_006799	Hs.72026	protease, serine, 21 (testisin)	1.34 2.19	2.19 2.79
	411789 411800	AF245505 N39342	Hs.72157 Hs.103042	Adlican microtubute-associated protein 18	23.34	34.00
_	411945	AL033527	Hs.92137	v-myc avian myelocytomatosis viral oncog	1.00	8.00
80	412115	AK001763	Hs.73239	hypothetical protein FLJ10901	2.07	1.64
	412140	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines	118.48	92.00
	412276	BE262621	Hs.73798	macrophage migration inhibitory factor (1.98	1.49
	412464	T78141	Hs.22826	ESTs, Weakly similar to I55214 salivary	1.16 41.52	1.34 84.00
85	412530 412537	AA766268 AL031778	Hs.266273	hypothetical protein FLJ13346 nuclear transcription factor Y, alpha	41.52 17.90	55.00
55	414001	1001110		timesom aminombront tonon, 12 mbito		

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	412659	AW753865	Hs.74376	olfactomedin related ER localized protei	14.65	47.00
	412719	AW016610	Hs.816	ESTs	382.46	128.00
	412723 412811	AA648459 H06382	Hs.335951	hypothetical protein AF301222 ESTs	54.90 1.00	1.00 11.00
5	412817	AL037159	Hs.74619	proteasome (prosome, macropain) 26S subu	1.63	1.42
_	412863	AA121673	Hs.59757	zinc finger protein 281	17.63	56.00
	412924	BE018422	Hs.75258	H2A histone family, member Y	1.00 2.19	22.00 2.05
	413004 413011	T35901 AW068115	Hs.75117 Hs.821	interleukin enhancer binding factor 2, 4 biglycan	1.22	1.88
10	413048	M93221	Hs.75182	mannose receptor, C type 1	0.30	6.23
	413063	AL035737	Hs.75184	chitinase 3-like 1 (cartilage glycoprote	3.43	8.71
	413129 413142	AF292100 M81740	Hs.104613 Hs.75212	RP42 homolog omithine decarboxylase 1	4.67 1.92	4.77 2.59
	413223	AI732182	Hs.191866	ESTs	5.73	27.00
15	413248	T64858	Hs.21433	hypothetical protein DKFZp547J038	0.99	1.06
	413273	U75679	Hs.75257	stern-loop (histone) binding protein	1.00 1.10	18.00 1.09
	413278 413281	BE563085 AA861271	Hs.833 Hs.222024	interferon-stimulated protein, 15 kDa transcription factor BMAL2	95.94	69.00
	413364	BE536218	Hs.137516	fidgetin-fike 1	1.00	1.00
20	413385	M34455	Hs.840	indoleamine-pyrrole 2,3 dioxygenase	0.95	2.09 1.00
	413409 413453	Al638418 AA129640	Hs.1440 Hs.128065	DEAD/H (Asp-Glu-Ala-Asp/His) box polypep ESTs	1.00 1.00	31.00
	413527	BE250788	Hs.179882	hypothetical protein FLJ12443	1.08	1.46
0.5	413554	AA319146	Hs.75426	secretogranin II (chromogranin C)	79.15	114.00
25	413573	AI733859	Hs.149089	ESTs	1.00 8.80	1.00 10.00
	413582 413597	AW295647 AW302885	Hs.71331 Hs.117183	hypothetical protein MGC5350 ESTs	1.00	1.00
	413690	BE157489	1.0.1111100	gb:RC1-HT0375-120200-011-e06 HT0375 Hamo	1.00	1.00
20	413691	AB023173	Hs.75478	ATPase, Class VI, type 11B	3.16	2.32
30	413719 413753	BE439580 U17760	Hs.75498 Hs.75517	small inducible cytokine subfamily A (Cy laminin, beta 3 (nicein (125kD), kalinin	2.88 144.10	9.52 108.00
	413801	M62246	Hs.35406	ESTs, Highly similar to unnamed protein	1.00	17.00
	413833	Z15005	Hs.75573	centromere protein E (312kD)	1.00	1.00
35	413882 413926	AA132973 AA133338	Hs.184492 Hs.54310	ESTs ESTs	64.24 1.00	148.00 67.00
55	413943	AW294416	Hs.144687	Homo sapiens cDNA FLJ12981 fis, clone NT	43.42	42.00
	413995	BE048146	Hs.75671	syntaxin 1A (brain)	1.23	1.11
	414035 414142	Y00630 AW368397	Hs.75716 Hs.334485	serine (or cystelne) proteinase inhibito Homo sapiens cDNA FLJ14438 fis, clone HE	2.02 1.00	2.51 102.00
40	414180	AIB63304	Hs.120905	Homo sapiens cDNA FLJ11448 fis, clone HE	6.92	77.00
	414245	BE148072	Hs.75850	WAS protein family, member 1	1.00	1.00
	414275	AW970254 BE263280	Hs.889 Hs.75888	Charot-Leyden crystal protein phosphogluconate dehydrogenase	1.00 1.52	59.00 1.73
	414317 414334	AA824298	Hs.21331	hypothetical protein FLJ10036	1.78	1.72
45	414341	D80004	Hs.75909	KIAA0182 protein	33.90	151.00
	414368 414416	W70171 AW409985	Hs.75939 Hs.76084	uridine monophosphate kinase hypothetical protein MGC2721	171.60 2.32	97.00 1.85
	414430	AI346201	Hs.76118	ubiquitin carboxyl-terminal esterase L1	226.15	66.00
50	414570	Y00285	Hs.76473	insulin-like growth factor 2 receptor	1.64	1.98
50	414618 414675	A1204600 R79015	Hs.96978 Hs.296281	hypothetical protein MGC10764 Interleukin enhancer binding factor 1	1.87 1.51	72.00 1.39
	414683	S78296	Hs.76888	hypothetical protein MGC12702	43.61	64.00
	414696	AF002020	Hs.76918	Niemann-Pick disease, type C1	28.63	71.00
55	414711 414718	Al310440 H95348	Hs.288735 Hs.107987	Homo sapiens cDNA FLJ13522 fis, clone PL ESTs	14.86 1.00	42.00 5.00
55	414732	AW410976	Hs.77152	minichromosome maintenance deficient (S.	1.64	1.44
	414747	U30872	Hs.77204	centromere protein F (350/400kD, mitosin	65.01	74.00
	414761	AU077228	Hs.77256	enhancer of zeste (Drosophila) homolog 2	130.35 2.24	121.00 2.19
60	414774 414806	X02419 D14694	Hs.77274 Hs.77329	plasminogen activator, urokinase phosphatidylserine synthase 1	1.63	1.53
-	414809	A1434699	Hs.77356	transferrin receptor (p90, CD71)	1.97	2.60
	414812	X72755	Hs.77367	monokine induced by gamma interferon	3.48 103.22	10.60 143.00
	414825 414839	X06370 X63692	Hs.77432 Hs.77462	epidermal growth factor receptor (avian DNA (cytosine-5-)-methyltransferase 1	1.80	1.69
65	414883	AA926960		CDC28 protein kinase 1	14.29	10.06
	414907	X90725	Hs.77597	poto (Drosophia)-like kinase	1.95 3.00	2.20 2.90
	414914 414945	U49844 BE076358	Hs.77613 Hs.77667	ataxia telangiectasia and Rad3 related lymphocyte antigen 6 complex, locus E	1.02	1.21
	414972	BE263782	Hs.77695	KIAA0008 gene product	1.00	1.00
70	415014	AW954064	Hs.24951	ESTs	1.42	2.84
	415091 415138	AL044872 C18356	Hs.77910 Hs.295944	3-hydroxy-3-methylglutaryl-Coenzyme A sy tissue factor pathway inhibitor 2	1.00 34.72	30.00 107.00
	415227	AW821113	Hs.72402	ESTs	1.87	49.00
75	415238	R37780	Hs.21422	ESTs	1.00	1.00
75	415263 415295	AA948033 R41450	Hs.130853 Hs.6546	ESTs ESTs	1.00 1.00	1.00 1.00
	415339	NM_015156	Hs.78398	KIAA0071 protein	51.18	166.00
	415669	NM_005025	Hs.78589	serine (or cysteine) proteinase inhibito	30.84	63.00
80	415674	BE394784	Hs.78596	proteasome (prosome, macropain) subunit,	1.48 1.00	1.39 1.00
υU	415709 415735	AA649850 AA704162	Hs.278558 Hs.120811	ESTs ESTs, Wealdy similar to 138022 hypotheti	1.00	72.00
	415799	AA653718	Hs.225841	DKFZP434D193 protein	6.23	31.00
	415817 415857	U88967 AA866115	Hs.78867 Hs.127797	protein tyrosine phosphatase, receptor-t Homo sapiens cDNA FLJ11381 fis, clone HE	24.30 32.51	1.00 35.00
85	415989	Al267700	110.121171	ESTs	78.89	1.00

	w	O 02/086	443			
	416018	AW138239	Hs.78977	proprotein convertase subtilisin/kexin t	1.00	1.00
	416065	BE267931	Hs.78996	proliferating cell nuclear antigen	3.35	2.32
	416111	AA033813	Hs.79018	chromatin assembly factor 1, subunit A (39.03	3.00
5	416177 416178	AA174069 AI808527	Hs.187607	ESTs serologically defined breast cancer anti	1.00 3.83	9.00 3.76
,	416208	AW291168	Hs.192822 Hs.41295	ESTs, Weakly similar to MUC2_HUMAN MUCIN	3.67	1.00
	416209	AA236776	Hs.79078	MAD2 (mitotic arrest deficient, yeast, h	9.70	1.00
	416239	AL038450	Hs.48948	ESTs	83.87	129.00
10	416250 416322	AA581386 BE019494	Hs.73452 Hs.79217	hypothetical protein MGC10791 pymoline-5-carboxylate reductase 1	1.96 2.08	2.12 1.73
10	416423	H54375	Hs.268921	ESTs	1.00	89.00
	416448	L13210	Hs.79339	lectin, galactoside-binding, soluble, 3	1.28	1.54
	416498	U33632	Hs.79351	potassium channel, subfamily K, member 1	27.29	67.00
15	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara	53.29 9.96	51.00 5.00
13	416661 416722	AA634543 AA354604	Hs.79440 Hs.122546	IGF-II mRNA-binding protein 3 hypothetical protein FLJ23017	3.68	33.00
	416819	U77735	Hs.80205	pim-2 oncogene	1.59	1.84
	416936	N21352	Hs.42987	ESTs, Weakly similar to S21348 probable	1.00	1.00
20	417034	NM_006183	Hs.80962	neurotensin	1.00 32.95	1.00 156.00
20	417061 417079	A1675944 U65590	Hs.188691 Hs.81134	Homo sapiens cDNA FLJ12033 fis, clone HE interleukin 1 receptor antagonist	3.91	4.93
	417218	AA129547	Hs.285754	met proto-oncogene (hepatocyte growth fa	1.00	51.00
	417233	W25005	Hs.24395	small inducible cytokine subfamily B (Cy	3.38	2.05
25	417308 417315	H60720	Hs.81892	KIAA0101 gene product ribosomal protein S24	82.94 106.61	25.36 121.00
23	417324	A1080042 AW265494	Hs.180450	ESTs	1.20	1.28
	417366	BE185289	Hs.1076	small proline-rich protein 18 (comifin)	8.97	3.27
	417389	BE260964	Hs.82045	midkine (neurite growth-promoting factor	2.59	1.82
30	417428	N87579	Hs.278871	gb:LL2030F Human fetal heart, Lambda ZAP	1.00 304.75	52.00 173.00
50	417433 417466	BE270266 Al681547	Hs.82128 Hs.59457	5T4 oncofetal trophoblast glycoprotein hypothetical protein FLJ22127	1.24	1.34
	417512	A1979168	Hs.344096	glycoprotein (transmembrane) nmb	2.14	5.50
	417515	L24203	Hs.82237	ataxia-telangiectasia group D-associated	2.66	1.68
35	417542	J04129	Hs.82269	progestagen-associated endometrial prote	1.28 42.76	1.35 51.00
22	417576 417715	AA339449 AW969587	Hs.82285 Hs.86366	phosphoribosylglycinamide formyltransfer ESTs	6.35	2.75
	417720	AA205625	Hs.208067	ESTs	113.31	56.00
	417791	AW965339	Hs.111471	ESTs	39.98	16.00
40	417830 417866	AW504786 AW067903	Hs.122579 Hs.82772	hypothetical protein FLJ 10461 collagen, type XI, alpha 1	2.61 2.35	31.00 2.44
40	417900	BE250127	Hs.82906	CDC20 (cell division cycle 20, S. cerevi	1.52	1.11
	417933	X02308	Hs.82962	thymidylate synthetase	4.74	2.55
	417944	AU077196	Hs.82985	collagen, type V, alpha 2	3.61	5.21
45	417975 417991	AA641836 AA731452	Hs.30085 Hs.190008	hypothetical protein FLJ23186 ESTs	12.49 1.00	38.00 26.00
15	418004	U37519	Hs.87539	aldehyde dehydrogenase 3 family, member	3.02	2.12
•	418007	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial	187.59	1.00
	418054	NM_002318	Hs.83354	lysyl oxidase-like 2	2.85 1.54	2.63 1.69
50	418057 418113	NM_012151 Al272141	Hs.83363 Hs.83484	coagulation factor VIII-associated (intr SRY (sex determining region Y)-box 4	6.82	5.22
	418140	BE613836 .	Hs.83551	microfibrillar-associated protein 2	- 1.26	1.46
	418203	X54942	Hs.83758	CDC28 protein kinase 2	134.19	144.00
	418207 418216	C14685 AA662240	Hs.34772 Hs.283099	ESTs AF15q14 protein	1.00 64.66	1.00 61.00
55	418236	AW994005	Hs.337534	ESTs	18.53	147.00
	418249	H89226	Hs.34892	KIAA1323 protein	30.53	106.00
	418281	U09550	Hs.1154	oviductal glycoprotein 1, 120kD (mucin 9	1.00	3.00
	418283 418300	S79895 Al433074	Hs.83942 Hs.86682	cathepsin K (pycnodysostosis) Homo sapiens cDNA: FLJ21578 fis, clone C	3.96 3.18	5.16 2.91
. 60	418322	AA284166	Hs.84113	cyclin-dependent kinase inhibitor 3 (CDK	11.96	6.68
	418327	U70370	Hs.84136	paired-like homeodomain transcription fa	9.23	2.22
	418345	AJ001696	Hs.241407	serine (or cysteine) proteinase inhibito	1.00 21.68	1.00 44.00
	418379 418397	AA218940 NM_001269	Hs.137516 Hs.84746	fidgetin-like 1 chromosome condensation 1	1.00	8.00
65	418403	D86978	Hs.84790	KIAA0225 protein	16.91	18.98
	418462	BE001596	Hs.85266	integrin, beta 4	1.56	1.16
	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me	3.22 2.66	2.38 2.22
	418506 418526	AA084248 BE019020	Hs.85339 Hs.85838	G protein-coupled receptor 39 solute carrier family 16 (monocarboxylic	2.04	2.21
70	418538	BE244323	Hs.85951	exportin, tRNA (nuclear export receptor	1.33	37.00
	418543	NM_005329	Hs.85962	hyaluronan synthase 3	1.04	1.23
	418574 418592	N28754 X99226	Uc 284153	M-phase phosphoprotein 9 Fanconi anemia, complementation group A	48.60 18.24	85.00 26.00
	418641	BE243136	Hs.284153 Hs.86947	a disintegrin and metalloproteinase doma	1.19	1.41
75	418661	NM_001949	Hs.1189	E2F transcription factor 3	29.05	43.00
	418663	AK001100	Hs.41690	desmocollin 3	112.17	19.00
	418678	NM_001327	Hs.87225	cancer/testis antigen	1.18 1.54	1.10 1.98
	418686 418689	Z36830 Al360883	Hs.87268 Hs.274448	ennexin A8 hypothetical protein FLJ 11029	1.19	1.04
80	418712	Z42183		gb:HSC0BF041 normalized infant brain cDN	1.00	12.00
	418727	AA227609	Hs.94834	ESTs	1.00	49.00
	418738 418819	AW388633 AA228776	Hs.6682 Hs.191721	solute carrier family 7, (calionic amino ESTs	49.85 1.00	1.00 140.00
0.5	418830	BE513731	Hs.68959	hypothetical protein MGC4816	20.97	23.00
85	418882	NM_004996	Hs.89433	ATP-binding cassette, sub-family C (CFTR	57.09	35.00

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	418971	AA360392	Hs.87113	ESTs	1.00	12.00
	418973	AA233056	Hs.191518	ESTs	4.89	28.00
	419078	M93119	Hs.89584	insulinoma-associated 1	1.00	10.00
_	419079	AW014836	Hs.18844	ESTs	1.09	1.98
5	419080	AW150835	Hs.18878	hypothetical protein FLJ21620	2.06	1.68 51.00
	419088	AI538323	Hs.52620	integrin, beta 8	15.60 1.11	1,83
	419092	J05581	Hs.89603	mucin 1, transmembrane parathyroid hormone-like hormone	1.00	1.00
	419121	AA374372	Hs.89626 Hs.89655	protein tyrosine phosphatase, receptor t	1.10	1.14
10	419171 419183	NM_002846 U60669	Hs.89663	cytochrome P450, subfamily XXIV (vitamin	1.00	1.00
10	419216	AU076718	Hs.164021	small inducible cytokine subfamily B (Cy	3.18	2.43
	419288	AA256106	Hs.87507	ESTs	1.00	34.00
	419335	AW960146	Hs.284137	hypothetical protein FLJ12888	1.00	8.00
	419354	M62839	Hs.1252	apolipoprotein H (beta-2-glycoprotein I)	22.63	54.00
15	419359	AL043202	Hs.90073	chromosome segregation 1 (yeast homolog)	2.50	1.98
	419423	D26488	Hs.90315	KIAA0007 protein	1.00	7.00
	419443	D62703		gb:HUM316G10B Clontech human aorta polyA	1.00	12.00
	419452	U33635	Hs.90572	PTK7 protein tyrosine kinase 7	1.64	1.84
00	419474	AW968619	Hs.155849	ESTs	13.63	62.00
20	419485	AA489023	Hs.99807	ESTs, Weakly similar to unnamed protein	4.27	2.26
	419488	AA316241	Hs.90691	nucleophosmin/nucleoplasmin 3	3.66	3.63
	419502	AU076704		fibrinogen, A alpha polypeptide	13.05	115.00
	419539	AF070590	Hs.90869	Homo sapiens clones 24622 and 24623 mRNA	74.60	117.00 4.98
25	419556	U29615	Hs.91093	chitinase 1 (chitotriosidase)	1.47 1.00	4.90
23	419569	AI971651	Hs.91143 Hs.91417	jagged 1 (Alagille syndrome) topolsomerase (DNA) Il binding protein	94.30	94.00
	419594 419703	AA013051 Al793257	Hs.128151	ESTs	15.26	50.00
	419721	NM_001650	Hs.288650	aquaporin 4	1.00	191.00
	419729	AA586442	Hs.21411	gb:no53a03.s1 NCI_CGAP_SS1 Homo sapiens	1.00	59.00
30	419741	NM_007019	Hs.93002	ubiquitin carrier protein E2-C	2.02	1.08
	419745	AF042001	Hs.93005	slug (chicken homolog), zinc finger prot	1.00	1.00
	419752	AA249573	Hs.152618	ESTs, Moderately similar to ZN91_HUMAN Z	29.87	77.00
	419839	U24577	Hs.93304	phospholipase A2, group VII (platelet-ac	50.99	214.00
	419936	Al792788		gb:ol91d05.y5 NCI_CGAP_Kid5 Homo sapiens	1.00	1.00
35	419937	AB040959	Hs.93836	DKFZP434N014 protein	1.64	2.47
	419983	W55956	Hs.94030	Homo sapiens mRNA; cDNA DKFZp586E1624 (f	15.72	94.00
	420005	AW271106	Hs.133294	ESTs	3.15	1.43
	420047	A1478658	Hs.94631	brefeldin A-inhibited guanine nucleotide	12.45 1.00	39.00 117.00
40	420058	AK001423	Hs.94694	Homo sapiens cDNA FLJ10561 fis, clone NT	1.43	1.21
40	420162 420251	BE378432 AW374968	Hs.95577 Hs.348112	cyclin-dependent kinase 4 Human DNA sequence from clone RP5-1103G7	2.35	3.23
	420259	AF004884	Hs.96253	calcium channel, voltage-depandent, P/Q	0.77	1.15
	420281	Al623693	Hs.323494	ESTs	45.04	54.00
	420309	AW043637	Hs.21766	ESTs, Weakly similar to ALU5_HUMAN ALU S	49.22	31.00
45	420332	NM_001756	Hs.1305	serine (or cystelne) proteinase inhibito	0.05	2.82
	420380	AA640891	Hs.102406	ESTs	0.99	2.74
	420462	AF050147	Hs.97932	chondromodulin I precursor	1.00	1.00
	420520	AK001978	Hs.98510	similar to rab11-binding protein	49.74	133.00
50	420552	AK000492	Hs.98806	hypothetical protein	94.65	88.00
50	420560	AW207748	Hs.59115	ESTs	1.00	17.00
*	420610	AI683183	Hs.99348	distal-less homeo box 5	1.00	13.00
	420689	H79979	Hs.88678	ESTs	50.09 1.00	95.00 31.00
	420721	AA927802	Hs.159471	ZAP3 protein Homo sapiens cDNA FLJ11381 fis, clone HE	1.00	48.00
55	420759 420783	T11832 Al659838	Hs.127797 Hs.99923	lectin, galactoside-binding, soluble, 7	3.04	1.25
55	420900	AL045633	Hs.44269	ESTs	2.24	7.00
	420931	AF044197	Hs.100431	small inducible cytokine B subfamily (Cy	1.00	8.00
	421002	AF116030	Hs.100932	transcription factor 17	1.00	27.00
	421027	AA761198	Hs.55254	ESTs	2.87	38.00
60	421037	AI684808	Hs.197653	ESTs	1.00	46.00
	421041	N36914	Hs.14691	ESTs, Moderately similar to 138022 hypot	1.00	98.00
•	421073	NM_004689	Hs.101448	metastasis associated 1	1.34	1.46
	421110	AJ250717	Hs.1355	cathepsin E	119.47	427.00
65	421133	AA401369	Hs.190721	ESTs	1.10	17.00 1.63
U.S	421150	Al913562	Hs.189902	ESTs	1.45 1.00	15.00
	421155 421307	H87879 BE539976	Hs.102267 Hs.103305	lysyl oxidase Homo sapiens mRNA; cDNA DKFZp434B0425 (f		1.10
	421316	AA287203	Hs.324728	SMA5	1.00	21.00
	421379	Y15221	Hs.103982	small inducible cytokine subfamily B (Cy	1.92	3.94
70	421451	AA291377	Hs.50831	ESTs	5.89	14.00
. •	421474	U76362	Hs.104637	solute carrier family 1 (glutamate trans	1.46	1.76
	421506	BE302796	Hs.105097	thymidine kinase 1, soluble	1.56	1.08
	421508	NM_004833	Hs.105115	absent in melanoma 2	5.11	5.23
~-	421515	Y11339	Hs.105352	GalNAc aipha-2, 6-sialyltransferase I, I	1.00	3.00
75	421524	AA312082	Hs.105445	GDNF family receptor alpha 1	2.63	10.58
	421526	AL080121	Hs.105460	DKFZP564O0823 protein	1.46	1.88
	421552	AF026692	Hs.105700	secreted frizzled-related protein 4	30.21	50.32
	421574	AJ000152	Hs.105924	defensin, beta 2	1.67 1.23	1.74 1.00
80	421582	AI910275	Un toegen	trefoil factor 1 (breast cancer, estroge	1.23 1.00	116.00
ou.	421633 421659	AF121860 NM_014459	Hs.106260 Hs.106511	sorting nexin 10 protocadherin 17	0.05	6.33
	421677	H64092	Hs.38282	ESTs	1.31	1.42
	421753	BE314828	Hs.107911	ATP-binding cassette, sub-family B (MDR/	1.41	1.20
	421773	W69233	Hs.112457	ESTs	1.12	1.14
85	421777	BE562088	Hs.108196	HSPC037 protein	1.97	1.29
				-		

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	421800	AA298151	Hs.222969	ESTs	1.03	1.30
	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	1.88	1.59
	421896	N62293	Hs.45107	ESTs .	11.84	22.80
5	421928	AF013758	Hs.109643	polyadenylate binding protein-Interactin gamma-aminobutyric acid (GABA) A recepto	45.89 1.13	90.00 1.49
,	421931 421948	NM_000814 L42583	Hs.1440 Hs.334309	keratin 6A	51.83	20.25
	421975	AW961017	Hs.6459	hypothetical protein FLJ11856	1.17	1.15
	422026	U80736	Hs.110826	trinucleotide repeat containing 9	1.00	52.00
10	422094	AF129535	Hs.272027	F-box only protein 5	67.61	62.00
10	422095	AI868872	Hs.282804	hypothetical protein FLJ22704	4.37	2.34
	422109	S73265	Hs.1473	gastrin-releasing peptide gb:QV0-OT0033-010400-182-a07 OT0033 Homo	4.18 40.89	95.50 71.00
	422128 422129	AW881145 AU076635	Hs.1478	serine (or cysteine) proteinase inhibito	1.13	1.38
	422134	AW179019	Hs.112110	mitochondrial ribosomal protein L42	41.59	96.00
15	422158	L10343	Hs.112341	protease Inhibitor 3, skin-derived (SKAL	2.37	1.10
	422168	AA586894	Hs.112408	S100 calcium-binding protein A7 (psorias	3.29	1.68
	422278	AF072873	Hs.114218	frizzled (Drosophila) homolog 6	4.93	5.73
	422282	AF019225	Hs.114309	apolipoprotein L	1.49 25.99	1.71 10.91
20	422283 422310	AW411307 AA316622	Hs.114311 Hs.98370	CDC45 (cell division cycle 45, S.cerevis cytochrome P450, subfamily IIS, polypept	1.54	1.41
20	422311	AF073515	Hs.114948	cytokine receptor-like factor 1	1.15	1.78
	422330	D30783	Hs.115263	epiregulin	1.00	112.00
	422364	AF067800	Hs.115515	C-type (calcium dependent, carbohydrate-	9.39	60.00
25	422406	AF025441	Hs.116206	Opa-interacting protein 5	18.33	53.00
25	422424	AI186431	Hs.296638	prostate differentiation factor aldo-keto reductase family 1, member B10	1.71 47.53	3.21 32.00
	422440 422487	NM_004812 AJ010901	Hs.116724 Hs.198267	mucin 4, tracheobronchial	73.68	35.54
	422511	AU076442	Hs.117938	collagen, type XVII, alpha 1	173.97	26.00
	422515	AW500470	Hs.117950	multifunctional polypeptide similar to S	4.68	2.92
30	422656	A1870435	Hs.1569	LIM homeobox protein 2	1.00	1.00
	422737	M26939	Hs.119571	collagen, type III, alpha 1 (Ehlers-Dani	3.89	4.55
•	422756 422765	AA441787	Hs.119689 Hs.1578	glycoprotein hormones, alpha polypeptide baculoviral IAP repeat-containing 5 (sur	1.05 3.88	1.46 1.53
	422809	AW409701 AK001379	Hs.121028	hypothetical protein FLJ 10549	99.56	53.00
35	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	1.69	3.17
	422938	NM_001809	Hs.1594	centromere protein A (17kD)	70.46	61.00
	422956	BE545072	Hs.122579	ECT2 protein (Epithelial cell transformi	77.74	3.00
	422960	AW890487	Hs.63984	cadherin 13, H-cadherin (heart)	5.88 171.41	8.55 17.00
40	422963 422976	AA401369 AU076657	Hs.190721 Hs.1600	ESTs chaperonin containing TCP1, subunit 5 (e	2.12	1.62
70	422981	AF026445	Hs.122752	TATA box binding protein (TBP)-associate	10.49	35.00
•	422986	AA319777	Hs.221974	ESTs	12.40	32.47
	423034	AL119930		gb:DKFZp761A092_r1 761 (synonym: hamy2)	16.41	60.00
15	423049	X59373	Hs.188023	ESTs, Moderately similar to HXDA_HUMAN H	1.00	1.00
45	423081	AF262992	Hs.123159	sperm associated antigen 4	1.82 1.14	2.96 1.53
	423184 423217	NM_004428 NM_000094	Hs.1624 Hs.1640	ephrin-A1 collagen, type VII, alpha 1 (epidermolys	2.14	1.69
	423248	AA380177	Hs.125845	ribulose-5-phosphate-3-epimerase	7.18	14.00
	423309	BE006775	Hs.126782	sushi-repeat protein	21.90	64.00
50	423361	AW170055	Hs.47628	ESTs	1.00	1.00
	423453	AW450737	Hs.128791	CGI-09 protein	55.52 0.88	66.00 1.17
	423511 423516	AF036329 AB007933	Hs.129715 Hs.129729	gonadotropin-releasing hormone 2 ligand of neuronal nitric oxide synthase	1.76	5.40
	423551	AA327598	Hs.233785	ESTs	3.54	4.33
55	423554	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	1.00	50.00
	423575	C18863	Hs.163443	Homo sapiens cDNA FLJ11576 fis, clone HE	38.88	70.00
	423624	AI807408	Hs.166368	ESTs	1.00	67.00 1.00
	423634 423642	AW959908 AW452650	Hs.1690 Hs.157148	heparin-binding growth factor binding pr hypothetical protein MGC13204	76.02 19.14	58.00
60	423662	AA642452	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro	3.61	13.57
	423673	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage	240.73	40.00
	423698	AA329796	Hs.1098	DKFZp434J1813 protein	1.00	59.00
	423725	AJ403108	Hs.132127	hypothetical protein LOC57822	4.20	1.00
65	423761 423787	NM_006194 AJ295745	Hs.132576 Hs.236204	paired box gene 9 nuclear pore complex protein	1.00 7.18	1.00 6.64
05	423816	AF151064	113.230204	hypothetical protein	1.00	44.00
	423826	U20325	Hs.1707	cocaine- and amphetamine-regulated trans	1.00	1.00
	423849	AL157425	Hs.133315	Homo saplens mRNA; cDNA DKFZp761J1324 (f	1.00	1.00
70	423887	AL080207	Hs.134585	DKFZP434G232 protein	1.00	1.00
70	423934	U89995	Hs.159234	forkhead box E1 (thyroid transcription f	31.33 5.81	31.00 10.87
	423954 423961	AW753164 D13666	Hs.288604 Hs.136348	KIAA1632 protein osteoblast specific factor 2 (fasciclin	3.55	3.30
	424012	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog	233.42	68.00
	424016	AW163729	Hs.6140	hypothetical protein MGC15730	0.93	1.01
75	424028	AF055084	Hs.153692	Homo sapiens cDNA FLJ14354 fis, clone Y7	21.30	52.00
	424046	AF027866	Hs.138202	serine (or cysteine) proteinase inhibito	1.00	1.00 70.00
	424086 424098	Al351010 AF077374	Hs.102267 Hs.139322	lysyl oxidase small proline-rich protein 3	21.91 137.82	54.00
	424120	T80579	Hs.290270	ESTs	1.00	1.00
80	424165	AW582904	Hs.142255	islet amyloid polypeptide	1.00	34.00
	424200	AA337221		gb:EST41944 Endometrial tumor Homo sapie	13.06	48.00
	424279	L29306	Hs.171814	tryptophan hydroxylase (tryptophan 5-mon	1.00	1.00
	424308 424326	AW975531 NM_014479	Hs.154443 Hs.145296	minichromosome maintenance deficient (S. disintegrin protease	164.58 53.72	87.00 302.00
85	424340	AA339036	Hs.7033	ESTs	0.88	1.15
				•		-

	W	O 02/086	443			
	424351	BE622117	Hs.145567	hypothetical protein	0.93	1.03
	424364	AW383226	Hs.201189	ESTs, Wealdy similar to G01763 atrophin-	7.02 95.55	3.24 92.00
	424381 424411	AA285249 NM_005209	Hs.146329 Hs.146549	protein kinase Chk2 crystallin, beta A2	1.63	3.25
5	424420	BE614743	Hs.146688	prostaglandin E synthase	1.63	1.33
	424441	X14850	Hs.147097	H2A histone family, member X	1.82	1.29
	424502	AF242388	Hs.149585	lengsin	1.00 1.02	1.00 2.24
	424503 424513	X06256 BE385864	Hs.149609 Hs.149894	Integrin, alpha 5 (fibronectin receptor, mitochondrial translational initiation f	1.00	17.00
10	424539	L02911	Hs.150402	Activin A receptor, type I (ACVR1) (ALK	32.46	108.00
	424568	AF005418	Hs.150595	cytochrome P450, subfamily XXVIA, polype	3.40	2.58
	424602	AK002055	Hs.151046	hypothetical protein FLJ11193	31.87	25.00 2.37
	424629 424645	M906 56 NM_014682	Hs.151393 Hs.151449	glutamate-cysteine ligase, catalytic sub KIAA0535 gene product	3.58 1.00	1.00
15	424687	J05070	Hs.151738	matrix metalloproteinase 9 (gelatinase B	2.12	2.23
	424717	AW992292	Hs.152213	wingless-type MMTV integration site fami	1.00	1.00
	424834	AK001432	Hs.153408	Homo sapiens cDNA FLJ10570 fis, clone NT	56.19	12.00
	424840	D79987	Hs.153479 Hs.153591	extra spindle poles, S. cerevisiae, homo	2.65 1.23	1.30 1.05
20	424867 424905	AI024860 NM_002497	Hs.153704	Not56 (D. melanogaster)-like protein NIMA (never in mitosis gene a)-related k	21.35	1.00
20	424979	D87989	Hs.154073	UDP-galactose transporter related	1.36	1.35
	424999	AW953120		gb:EST365190 MAGE resequences, MAGB Homo	1.24	1.41
	425048	H05468	Hs.164502	ESTS	1.00 7.46	11.00 87.00
25	425057 425081	AA826434 X74794	Hs.1619 Hs.154443	achaete-scute complex (Drosophila) homol minichromosome maintenance deficient (S.	2.52	3.82
25	425118	AU076611	Hs.154672	methylene tetrahydrofolate dehydrogenase	4.84	4.03
•	425159	NM_004341	Hs.154868	carbamoyl-phosphate synthetase 2, aspart	3.62	2.73
	425202	AW962282	Hs.152049	ESTs, Weakly similar to 138022 hypotheti	1.00 100.77	53.00 44.00
30	425234 425236	AW152225 AW067800	Hs.165909 Hs.155223	ESTs, Weakly similar to 138022 hypotheti stanniocalcin 2	3.30	2.90
50	425245	A1751768	Hs.155314	KIAA0095 gene product	1.91	2.32
	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin	1.41	1.49
	425266	J00077	Hs.155421	alpha-fetoprotein	1.00	68.00
35	425274	BE281191	Hs.155462	minichromosome maintenance deficient (mi protein kinase, DNA-activated, catalytic	1.97 141.49	1.63 123.00
33	425322 425349	U63630 AA425234	Hs.155637 Hs.79886	ribose 5-phosphate isomerase A (ribose 5	1.00	84.00
	425371	D49441	Hs.155981	mesothelin	0.87	1.59
	425397	J04088	Hs.156346	topoisomerase (DNA) II alpha (170kD)	14.90	5.76
40	425420	BE536911	Hs.234545	hypothetical protein NUF2R	1.00 10.58	1.00 9.74
40	425424 425483	NM_004954 AF231022	Hs.157199 Hs.158159	ELKL motif kinase FAT turnor suppressor (Drosophila) homolo	1.74	1.40
		AW162943	Hs.250618	UL16 binding protein 2	1.49	1.14
	425580	L11144	Hs.1907	galanin	53.29	233.00
15	425650	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	33.45	1.00 55.00
45	425692 425695	D90041 NM_005401	Hs.155956 Hs.159238	N-acetyltransferase 1 (arylamine N-acety protein tyrosine phosphatase, non-recept	1.00 1.00	10.00
	425734	AF056209	Hs.159396	peptidylglycine alpha-amidating monooxyg	1.00	41.00
	425776	U25128	Hs.159499	parathyroid hormone receptor 2	1.00	48.00
50	425810	A1923627	Hs.31903	ESTs	27.39 1.99	98.00 1.58
30	425811 425849	AL039104 Al077288	Hs.159557 Hs.296323	karyopherin alpha 2 (RAG cohort 1, impor serum/glucocorticoid regulated kinase	71.16	3.42
	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily member	1.35	1.34
	426067	AA401369	Hs.190721	ESTs	1.01	17.00
55	426088	AF038007	Hs.166196	ATPase, Class I, type 8B, member 1	26.26 1.91	47.00 2.90
33	426215	AW067800	Hs.155223 Hs.154299	stannlocatcin 2 Human proteinase activated receptor-2 mR	22.40	25.00
	426227 426269	U67058 H15302	Hs.168950	Homo sapiens mRNA; cDNA DKFZp566A1046 (f	1.00	1.00
	426283	NM_003937	Hs.169139	kynureninase (L-kynurenine hydrolase)	91.39	229.00
60	426329	AL389951	Hs.271623	nucleoporin 50kD	4.34	4.08 1.00
60	426427 426432	M86699 AF001601	Hs.169840 Hs.169857	TTK protein kinase paraoxonase 2	7.02 1.16	1.68
	426440	BE382756	Hs.169902	solute carrier family 2 (facilitated glu	2.59	1.71
	426459	AF151812	Hs.169992	hypothetical 43.2 Kd protein	1.56	1.66
65	426471	M22440	Hs.170009	transforming growth factor, alpha	20.60	26.00
65	426496	D31765 AA401369	Hs.170114 Hs.190721	KIAA0061 protein ESTs	. 9.81 19.23	22.00 17.00
	426501 426514	BE616633	Hs.170195	bone morphogenetic protein 7 (osteogenic	103.74	41.00
	426536	A1949749	Hs.44441	ESTs	4.65	23.00
70	426572	AB037783	Hs.170623	hypothetical protein FLJ11183	1.00	43.00
70	426682	AV660038 NM_006201	Hs.2056	UDP glycosyltransferase 1 family, polype	160.06 1.51	8.00 1.35
	426691 426746	J03626	Hs.171834 Hs.2057	PCTAIRE protein kinase 1 uridine monophosphate synthetase (orotat	2.13	1.68
	426752	X69490	Hs.172004	tiin	0.02	5.14
75	426784	U03749	Hs.172216	chromogranin A (parathyroid secretory pr	1.72	1.71
75	426807	AA385315	Hs.156682	ESTs	1.30 1.47	1.64 1.53
	426812 426814	AF105365 AF036943	Hs.172613 Hs.172619	sclute carrier family 12 (potassium/chlo myelin transcription factor 1-like	1.00	1.00
	426831	BE296216	Hs.172673	S-adenosylhomocysteine hydrofase	1.51	1.25
0.0	426897	AA401369	Hs.190721	ESTs	141.56	17.00
80	426925	NM_001196	Hs.315689	Homo saplens cDNA: FLJ22373 fis, clone H	32.61	38.00
	426935 426964	NM_000088 AA393739	Hs.172928 Hs.287416	collagen, type I, alpha 1 Homo sapiens cDNA FLJ11439 fis, clone HE	2.65 1.97	3.16 3.49
	426966	Al493134	110.207410	sclerostin	1.00	1.00
0.5	426991	AK001536		Homo sapiens cDNA FLJ10674 fis, clone NT	3.39	2.28
85	427099	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophila, mous	4.24	17.00

	**/	A 02/096	443			
	427239	O 02/086 BE270447	443 Hs.174070	ubiquitin carrier protein	1.58	1.05
	427260	AA663848		gb:ae70b06.s1 Stratagene schizo brain S1	1.34	1.60
	427281	AA906147	Hs.102869	EST ₈	1.00	66.00 4.00
5	427335 427354	AA448542 T57896	Hs.251677 Hs.191095	G antigen 7B ESTs	51.83 1.17	1.95
,	427356	AW023482	Hs.97849	ESTs	7.31	41.00
	427376	AA401533	Hs.19440	ESTs	1.00	57.00
	427383	NM_005411	Hs.177582 Hs.177938	surfactant, pulmonary-associated protein lectin, superfamily member 1 (cartilage-	0.42 1.00	1.32 20.00
10	427427 427441	AF077345 AA412605	Hs.343879	SPANX family, member C	1.00	1.00
	427445	X80818	Hs.178078	glutamate receptor, metabotropic 4	0.97	1.03
	427505	AA361562	Hs.178761	26S proteasome-associated pad1 homolog	4.60	4.04 45.00
	427510 427528	Z47542 AU077143	Hs.179312 Hs.179565	small nuclear RNA activating complex, po minichromosome maintenance deficient (S.	22.00 97.45	92.00
15	427546	AA188763	Hs.36793	hypothetical protein FLJ23188	1.50	3.24
	427562	R56424	Hs.26534	ESTs	6.81	40.00
	427585	D31152	Hs.179729	collagen, type X, alpha 1 (Schmid metaph	69.91 2.70	62.00 49.00
	427660 427666	Al741320 Al791495	Hs.114121 Hs.180142	Homo sapiens cDNA: FLJ23228 fis, clone C calmodulin-like skin protein	1.37	1.88
20	427668	AA298760	Hs.180191	hypothetical protein FLJ14904	29.55	67.00
	427677	NM_007045	Hs.180296	FGFR1 oncogene partner	3.52	2.63
	427701	AA411101	Hs.243886	nuclear autoantigenic sperm protein (his	7.41 15.84	34.00 70.00
	427711 427719	M31659 Al393122	Hs.180408 Hs.134726	solute carrier family 25 (mitochondrial ESTs	7.03	4.52
25	427722	AK000123	Hs.180479	hypothetical protein FLJ20116	2.92	1.74
	427747	AW411425	Hs.180655	serine/threonine kinase 12	1.76	1.26
	427912	AL022310	Hs.181097	tumor necrosis factor (ligand) superfami	9.63 41.97	59.00 118.00
	427961 428004	AW293165 AA449563	Hs.143134 Hs.151393	ESTs glutamate-cysteine ligase, catalylic sub	23.82	1.00
30	428023	AL038843	1.0.101000	Homo sapiens cDNA: FLJ23602 fis, clone L	1.40	1.33
	428046	AW812795	Hs.337534	ESTs, Moderately similar to I38022 hypot	96.28	167.00
	428093	AW594506	Hs.104830	ESTs	1 <i>.2</i> 5 1.86	1.29 1.60
	428098 428129	AU077258 Al244311	Hs.182429 Hs.26912	protein disulfide isomerase-related prot ESTs	1.00	42.00
· 35	428169	Al928984	Hs.182793	golgi phosphoprotein 2	2.76	2.11
	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT	1.00	1.00
	428227	AA321649	Hs.2248 Hs.2250	small inducible cytokine subfamily B (Cy leukemia inhibitory factor (cholinergic	85.59 8.57	181.00 21.64
	428242 428330	H55709 L22524	Hs.2256	matrix metalloproteinase 7 (matrilysin,	7.77	15.90
40	428434	Al909935	Hs.65551	Homo saplens, Similar to DNA segment, Ch	0.58	1.43
	428450	NM_014791	Hs.184339	KIAA0175 gene product	237.53 6.00	204.00 4.60
	428471 428479	X57348 Y00272	Hs.184510 Hs.334562	stratifin cell division cycle 2, G1 to S and G2 to	56.54	16.00
	428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino	3.53	2.15
45	428505	AL035461	Hs.2281	chromogranin B (secretogranin 1)	1.00	1.00
	428532	AF157326	Hs.184786	TBP-interacting protein	1.00 1.00	58.00 16.00
	428645 428664	AA431400 AK001666	Hs.98729 Hs.189095	ESTs, Weakly similar to 2017205A dihydro similar to SALL1 (sal (Drosophila)-like	1.00	1.00
	428698	AA852773	Hs.334838	KIAA1866 protein	187.37	255.00
50	428728	NM_016625	Hs.191381	hypothetical protein	47.24	80.00
	428748 428758	AW593206 AA433988	Hs.98785 Hs.98502	Ksp37 protein hypothetical protein FLJ14303	1.00 1.06	87.00 1.13
	428771	AB028992	Hs.193143	KIAA1069 protein	1.98	92.00
	428801	AW277121	Hs.254881	ESTs	1.67	6.15
55	428810	AF068236	Hs.193788	nitric oxide synthase 2A (inducible, hep Homo sapiens cDNA FLJ14814 fis, clone NT	1.03 124.17	1.27 43.00
	428839 428845	. A1767756 AL157579	Hs.82302 Hs.153610	KIAA0751 gene product	1.00	1.00
	428959	AF100779	Hs.194680	WNT1 inducible signaling pathway protein	15.16	27.00
<u>(0</u>	428969	AF120274	Hs.194689	artemin	1.36	1.24
60	429038 429065	AL023513	Hs.194766 Hs.29643	seizure related gene 6 (mouse)-like Homo sapiens cDNA FLJ 13103 fis, clone NT	0.97 6.82	3.31 16.47
	429164	AI753247 AI688663	Hs.116586	ESTs	19.08	67.00
	429170	NM_001394	Hs.2359	dual specificity phosphatase 4	16.18	105.00
65	429183	AB014604	Hs.197955	KIAA0704 protein	79.72 1.00	104.00 1.00
05	429201 429211	X03178 AF052693	Hs.198246 Hs.198249	group-specific component (vitamin D bind gap junction protein, bela 5 (connexin 3	1.33	1.09
	429220	AW207206	7101,1004.10	ESTs	1.00	7.00
	429228	AI553633	Hs.326447	ESTs	39.47	29.25
70	429259	AA420450	Hs.292911	ESTs, Highly similar to S60712 band-6-pr ATP-binding cassette, sub-family A (ABC1	2.01 1.07	1.18 1.00
70	429263 429276	AA019004 AF056085	Hs.198396 Hs.198612	G protein-coupled receptor 51	3.70	142.00
	429359	W00482	Hs.2399	matrix metalloproteinase 14 (membrane-in	1.30	1.94
	429412	NM_006235	Hs.2407	POU domain, class 2, associating factor	94.09	86.00
75	429413 429486	NM_014058 AF155827	Hs.201877 Hs.203963	DESC1 protein hypothetical protein FLJ10339	41.91 12.19	10.00 1.00
, 5	429504	X99133	Hs.204238	lipocalin 2 (oncogene 24p3)	1.61	1.08
•	429538	BE182592	Hs.11261	small proline-rich protein 2A	4.43	2.90
	429547	AA401369	Hs.190721	ESTs	1.06 2.89	17.00 65.00
80	429551 429563	AW450624 BE619413	Hs.220931 Hs.2437	ESTs eukaryotic translation initiation factor	1.49	1.37
50	429597	NM_003816	Hs.2442	a disintegrin and matalloproteinase doma	61.86	100.00
	429610	AB024937	Hs.211092	LUNX protein; PLUNC (palate lung and nas	1.59	1.69
	429612 429616	AF062649 AI982722	Hs.252587	pituitary tumor-transforming 1 ESTs	2.78 1.00	1.74 1.00
85	429656	X05608	Hs.120845 Hs.211584	neurofilament, light polypeptide (68kD)	1.00	4.00

		O 02/086		and the same Pro-	60.05	104.00
	429663	M68874	Hs.211587	phospholipase A2, group IVA (cytosolic,	69.95 1.25	104.00 1.21
	429736 429782	AF125304 NM_005754	Hs.212680 Hs.220689	tumor necrosis factor receptor superfamil Ras-GTPase-activating protein SH3-domain	1.00	7.00
	429903	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	44.00	1.00
5	429918	AW873986	Hs.119383	ESTs	1.00	78.00
	429978	AA249027		ribosomal protein S6	1.98	3.09
	429986	AF092047	Hs.227277	sine oculis homeobox (Drosophila) homolo	1.00 69.27	48.00 59.00
	430044 430114	AA464510 AA847744	Hs.152812 Hs.99640	ESTs ESTs	1.00	1.00
10	430134	BE380149	Hs.105223	ESTs, Weakly similar to T33188 hypotheti	1.00	51.00
	430147	R60704	Hs.234434	hairy/enhancer-of-split related with YRP	1.10	2.22
	430287	AW182459	Hs.125759	ESTs, Weakly similar to LEU5_HUMAN LEUKE	1.00	127.00
	430294	AI538226	Hs.32976	guanine nucleotide binding protein 4	3.80	1.47
15	430300	U60805	Hs.238648	oncostatin M receptor	1.00 92,31	35.00 28.00
13	430315 430337	NM_004293 M36707	Hs.239147 Hs.239600	guanine deaminase calmodulin-like 3	1.18	1.08
	430378	Z29572	Hs.2556	tumor necrosis factor receptor superfami	5.28	66.00
	430388	AA356923	Hs.240770	nuclear cap binding protein subunit 2, 2	16.76	38.00
00	430393	BE185030	Hs.241305	estrogen-responsive B box protein	1.63	1.50
20	430439	AL133561		DKFZP434B061 protein	1.00	1.00 2.12
	430451	AA836472	Hs.297939	cathepsin B	1.64 63.35	44.00
	430454 430466	AW469011 AF052573	Hs.105635 Hs.241517	ESTs polymerase (DNA directed), theta	2.47	1.91
	430481	AA479678	Hs.203269	ESTs, Moderately similar to ALU8_HUMAN A	1.00	31.00
25	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam	12.28	41.00
	430508	Al015435	Hs.104637	ESTs	4.75	7.27
	430533	AA480895	Hs.57749	ESTs, Weakly similar to T17288 hypotheti	1.00	1.00
	430563	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	1.00 1.72	1.59 1.30
30	430677 430678	Z26317 AA401369	Hs.94560 Hs.190721	desmoglein 2 ESTs	0.90	17.00
50	430686	NM_001942	Hs.2633	desmoglein 1	1.00	1.00
	430788	Al742925	Hs.7179	ESTs, Weakly similar to 2004399A chromos	1.62	1.84
•	430890	X54232	Hs.2699	glypican 1	1.58	1.40
25	430935	AW072916		zinc finger protein 131 (clone pHZ-10)	90.28	132.00
35	430985	AA490232	Hs.27323	ESTs, Weakly similar to 178885 serine/th	0.94 60.25	1.28 28.00
•	431009 431089	BE149762 BE041395	Hs.48956	gap junction protein, beta 6 (connexin 3 ESTs, Weakly similar to unknown protein	23.32	941.00
	431092	Al332764	Hs.125757	ESTs	13.46	63.00
	431124	AF284221	Hs.59506	doublesex and mab-3 related transcriptio	49.43	62.00
40	431164	AA493650	Hs.94367	Homo sapiens cDNA: FLJ23494 fis, clone L	0.44	2.20
	431211	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	182.26	101.00
	431221	AW207837	Hs.286145	SRB7 (suppressor of RNA polymerase B, ye	4.15 1.00	13,97 86.00
	431277 431322	AA501806 AW970622	Hs.345824	ests gb:EST382704 MAGE resequences, MAGK Homo	40.55	200.00
45	431342	AW971018	Hs.21659	ESTs	1.00	53.00
	431384	BE158000	Hs.285026	gb:MR2-HT0377-150200-202-e03 HT0377 Homo	0.94	1.14
	431462	AW583672	Hs.256311	granin-like neuroendocrine peptide precu	1.30	1.25
	431494	AA991355	Hs.298312	hypothetical protein DKFZp434A1315	3.90 1.41	26.00 1.87
50	431515 431548	NM_012152 Al834273	Hs.258583 Hs.9711	endothetial differentiation, lysophospha novel protein	5.66	15.00
50	431630	NM_002204	Hs.265829	integrin, alpha 3 (antigen CD49C, alpha	0.99	1.44
	431745	AW972448	Hs.163425	ESTs	0.99	3.51
	431770	BE221880	Hs.268555	5'-3' exoribonuclease 2	67.12	91.00
E E	431830	Y16645	Hs.271387	small inducible cytokine subfamily A (Cy	3.36	4.71 2.51
55	431846	BE019924	Hs.271580	uroplakin 1B integrin, alpha 2 (CD49B, alpha 2 subuni	4.49 2.20	3.32
	431890 431934	X17033 AB031481	Hs.271986 Hs.272214	STG protein	1.01	1.04
	431958	X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta	51.17	46.35
	432006	AL137382	Hs.272320	Homo sapiens mRNA; cDNA DKFZp434L1226 (f	0.94	1.65
60 .	432023	R43020	Hs.236223	EST	0.94	47.00
	432201	Al538613	Hs.298241	Transmembrane protease, serine 3	1.10 1.42	2.24 1.45
	432210	AI567421	Hs.273330	Homo sapiens, clone IMAGE:3544662, mRNA, phosphate cytidylyltransferase 1, cholin	1.00	1.00
	432226 432239	AW182766 X81334	Hs.273558 Hs.2936	matrix metalloproteinase 13 (collagenase	18.67	1.00
65	432265	BE382679	Hs.285753	SCG10-like-protein	- 1.09	1.21
	432281	AK001239	Hs.274263	hypothetical protein FLJ10377	40.98	58.00
	432365	AK001106	Hs.274419	hypothetical protein FLJ10244	1.00	214.00
	432374	W68815	Hs.301885	Homo sapiens cDNA FLJ11346 fis, clone PL	157.34 1.65	37.00 1.06
70	432375	BE536069 AA221036	Hs.2962	S100 calcium-binding protein P gb:zr03f12.r1 Stratagene NT2 neuronal pr	73.71	75.00
70	432407 432441	AW292425	Hs.163484	ESTs	56.35	72.00
	432489	A1804855	Hs.207530	ESTs	1.00	24.00
	432543	AA552690	Hs.152423	Homo sapiens cDNA: FLJ21274 fis, clone C	137.72	98.00
75	432552	Al537170	Hs.173725	ESTs, Weakly similar to ALU8_HUMAN ALU S	1.00	31.00
75	432583	AW023624	Hs.162282	potassium channel TASK-4; potassium chan	0.27 2.87	35.18 6.22
	432606 432625	NM_002104 Al243596	Hs.3066 Hs.94830	granzyme K (serine protease, granzyme 3; ESTs, Moderately similar to T03094 A-kin	26.63	56.00
	432653	N62096	Hs.293185	ESTs, Weakly similar to JC7328 amino aci	1.92	5.29
	432677	NM_004482	Hs.278611	UDP-N-acetyl-alpha-D-galactosamine:polyp	1.00	48.00
80	432715	AA247152	Hs.200483	ESTs, Weakly similar to KIAA1074 protein	45.13	31.00
	432753	NM_014075	Hs.336938	Homo saplens PRO0593 mRNA, complete cds	1.00	68.00
	432788	AA521091	Hs.178499	Homo sapiens cDNA: FLJ23117 fis, clone L hypothetical protein MGC4485	2.69 1.22	3.67 1.34
	432842 432867	AW674093 AW016936	Hs.334822 Hs.233364	ESTs	1.00	1.00
85	432917	NM_014125	Hs.241517	PRO0327 protein	10.25	6.62
				•		

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	432920	U37689	Hs.3128	polymerase (RNA) II (DNA directed) polyp	1.44	1.30
	433001 433023	AF217513 AW864793	Hs.279905 Hs.87409	clone HQ0310 PRO0310p1 thrombospondin 1	154.79 20.96	85.64 100.00
	433042	AW193534	Hs.281895	Homo sepiens cDNA FLJ11660 fis, clone HE	1.00	10.00
5	433091	Y12642	Hs.3185	lymphocyte antigen 6 complex, locus D	1.20	1.09
	433159	AB035898	Hs.150587	kinesin-like protein 2	13.82	39.00
	433183	AF231338	Hs.222024	transcription factor BMAL2	1.00 1.00	69.00
	433258 433409	AA622788 Al278802	Hs.203613 Hs.25661	ESTs, Weakly similar to ALUB_HUMAN !!!! ESTs	44.81	1.25 117.00
10	433437	U20536	Hs.3280	caspase 6, apoptosis-related cysteine pr	70.39	105.00
- 0	433485	Al493076	Hs.201967	aldo-keto reductase family 1, member C2	11.55	2.00
	433537	AI733692	Hs.112488	ESTs	8.66	55.00
	433547	W04978	Hs.303023	beta tubulin 1, class VI	25.16	83.00
15	433556 433647	W56321	Hs.111460	calcium/calmodulin-dependent protein kin ESTs	1.00 20.30	19.00 49.00
13	433658	AA603367 L03678	Hs.222294 Hs.156110	immunoglobulin kappa constant	5.92	10.03
	433800	AI094221	Hs.135150	lung type-I ceil membrane-associated gly	2.29	2.22
	433819	AW511097	Hs.112765	ESTs	3.71	8.00
20	433862	D86960	Hs.3610	KIAA0205 gene product	62.08	104.00
20	433980	AA137152	Hs.286049	phosphoserine aminotransferase	108.91	47.00 1.00
	434088	AF116677 AA305599	Hs.249270 Hs.238205	hypothetical protein PRO1986 hypothetical protein PRO2013	1.00 121.27	87.00
	434094 434105	AW952124	Hs.13094	presenilins associated rhomboid-like pro	1.22	1.23
	434217	AW014795	Hs.23349	ESTs	14.11	57.00
25	434340	Al193043	Hs.128685	ESTs, Wealdy similar to T17226 hypotheti	2.10	2.56
	434360	AA401369	Hs.190721	ESTs	40.98	17.00
	434414	A1798376		gb:tr34b07.x1 NCI_CGAP_Ov23 Homo sapiens	1.48	1.56
	434424	Ai811202	Hs.325335	Homo sapiens cDNA: FLJ23523 fis, clone L	1.00 54.91	64.00 85.00
30	434467 434551	BE552368 BE387162	Hs.231853 Hs.280858	Homo sapiens cDNA FLJ 13445 fis, clone PL ESTs, Highly similar to A35661 DNA excis	2.46	2.00
50	434627	Al221894	Hs.39311	ESTs .	1.00	1.00
	434699	AA643687	Hs.149425	Homo sapiens cDNA FLJ11980 fis, clone HE	1.00	23.00
	434769	AA648884	Hs.134278	Homo sapiens cDNA FLJ12676 fis, clone NT	7.08	56.00
25	434792	AA649253	Hs.132458	ESTs	8.52	44.00
35	434808	AF155108	Hs.256150	Homo sapiens, Similar to RIKEN cDNA 2810	11.33 1.00	1.00 1.00
	434828 434876	D90070 AF160477	Hs.96 Hs.61460	phorbol-12-myristate-13-acetate-induced Ig superfamily receptor LNIR	1.25	1.29
	434891	AA814309	Hs.123583	ESTs	1.00	6.00
	434928	AW015595	Hs.4267	Homo sapiens clones 24714 and 24715 mRNA	1.00	1.00
40	435013	H91923	Hs.110024	Target CAT	1.26	1.10
	435066	BE261750	Hs.4747	dyskeratosis congenita 1, dyskerin	1.69	1.37
	435087	AW975241	Hs.23567	ESTS	1.00 2.90	1.00 1.93
	435099 435159	AC004770 AA668879	Hs.4756 Hs.116649	flap structure-specific endonuclease 1 ESTs	1.00	1.00
45	435205	X54136	Hs.181125	immunoglobulin tambda locus	1.02	1.46
15	435232	NM_001262	Hs.4854	cyclin-dependent kinase inhibitor 2C (p1	2.04	2.70
	435304	H10709	Hs.269524	ÉSTs	27.58	139.00
	435313	A1769400	Hs.189729	ESTs	1.00	14.00
50	435505	AF200492	Hs.211238	interleukin-1 homolog 1	1.00	38.00
50	435509	AM58679	Hs.181915 Hs.123310	ESTs ESTs	1.00 1.00	1.00 56.00
	435525 435532	Al831297 AW291488	Hs.117305	Homo sapiens, clone IMAGE:3682908, mRNA	1.00	2.00
	435550	A1224456	Hs.324507	H.sapiens polyA site DNA	3.42	3.92
	435602	AF217515	Hs.283532	uncharacterized bone marrow protein BM03	3.95	1.80
55	435766	R11673	Hs.186498	ESTs	1.00	28.00
	435793	AB037734	Hs.4993	KIAA1313 protein	23.68	42.00 58.00
	436069 436170	A1056879	Hs.263209 Hs.14529	ESTs .	1.00 1.00	18.00
	436211	AW450381 AK001581	Hs.334828	hypothetical protein FLI 10719; KIAA1794	5.84	22.00
60	436213	AA325512	Hs.71472	hypothetical protein FLJ10774; KIAA1709	1.42	1.27
	436217	T53925	Hs.107	fibrinogen-like 1	57.97	31.00
	436238	AK002163	Hs.301724	hypothetical protein FLJ11301	2.51	1.71
	436251	BE515065	Hs.296585	nucleolar protein (KKE/D repeat)	2.33	1.64
65	436291	BE568452 AL355841	Hs.344037 Hs.99330	protein regulator of cytokinesis 1 hypothetical protein FLJ23588	108.99 0.75	52.00 2.81
UJ	436302 436396	AU355641 AW992292	Hs.152213	wingless-type MMTV integration site fami	60.01	1.00
	436414	BE264633	Hs.143638	WD repeat domain 4	2.50	2.19
	436419	AI948626	Hs.171356	ESTs	0.95	1.33
70	436443	AW138211	Hs.128746	ESTs	1.12	9.26
70	436474	AJ270693	Hs.199887	ESTs	1.00	1.00
	436481 - 436486	AA379597 AA742221	Hs.5199 Hs.120633	HSPC150 protein similar to ubiquitin-con	3.28 1.00	1.56 19.00
	436511	AA721252	Hs.291502	ESTs ESTs	16.76	14.00
	436553	X57809	Hs.181125	immunoglobulin lambda locus	1.08	1.74
75	436557	W15573	Hs.5027	ESTs, Weakly similar to A47582 B-cell gr	19.20	9.75
	436608	AA628980		down syndrome critical region protein DS	33.92	25.00
	436667	AW025183	Hs.127680	ESTs	0.89	1.19
	436771	AW975687	Hs.292979	ESTS	1.00 1.00	10.00 17.00
80	436839 436887	AA401369 AW953157	Hs.190721 Hs.193235	ESTs hypothetical protein DKFZp547D155	1.06	1.15
55	436944	AW268614	Hs.5840	ESTs	1.00	1.00
	436961	AW375974	Hs.156704	ESTs	25.13	25.00
	436972	AA284679	Hs.25640	claudin 3	1.59	1.46
95	437016	AU076916	Hs.5398	guanine monphosphate synthelase	2.35	1.78
85	437044	AL035864	Hs.69517	cDNA for differentially expressed CO16 g	1.34	1.13

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	437181	Al305615	Hs.125343	ESTs, Weakly similar to KIAA0758 protein	1.00	17.00
	437204	AL110216	Hs.22826	ESTs, Weakly similar to 155214 safivery	40.55	82.00
	437205	AL110232	Hs.279243	Homo sapiens mRNA; cDNA DKFZp564D2071 (f	1.00	112.00
_	437259	Al377755	Hs.120695	ESTs	1.00	205.00
5	437270	R18087	Hs.323769	cisplatin resistance related protein CRR	1.56	1.54
	437271	AL137445	Hs.28846	Homo sapiens mRNA; cDNA DKFZp566O134 (fr	113.25	125.00
	437370	AL359567	Hs.161962	Homo sapiens mRNA; cDNA DKFZp547D023 (fr	1.82 1.35	4.57 1.75
	437390 437412	A1125859 BE069288	Hs.112607 Hs.34744	ESTs Homo sapiens mRNA; cDNA DKFZp547C136 (fr	3.58	3.20
10	437435	Al306152	Hs.27027	hypothetical protein DKFZp762H1311	3.03	1.08
10	437444	H46008	Hs.31518	ESTs	1.00	39.00
	437568	AJ954795	Hs.156135	ESTs	1.00	19.00
	437623	D63880	Hs.5719	chromosome condensation-related SMC-asso	1.95	1.57
1.5	437789	Al581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti	1.00	3.00
15	437814	AJ088192	Hs.135474	ESTs, Weakly similar to DDX9_HUMAN ATP-D	1.00	45.00
	437840	AA884836	Hs.292014	ESTs	1.07	1.78
	437852	BE001836	Hs.256897	ESTs, Weakly similar to dJ365012.1 [H.sa	1.68 1.87	3.26 2.52
	437879 437915	BE262082 Al637993	Hs.5894 Hs.202312	hypothetical protein FLJ10305 Homo sapiens clone N11 NTera2D1 teratoca	74.05	35.00
20	437916	BE566249	Hs.20999	hypothetical protein FLJ23142	23.15	89.00
20	437937	Al917222	Hs.121655	ESTs	1.00	1.00
	437942	Al888256	Hs.307526	ESTs	12.28	31.00
	438091	AW373062		nuclear receptor subfamily 1, group 1, m	1.53	10.85
0.5	438113	AJ467908	Hs.8882	ESTs	1.80	2.39
25	438119	AW963217	Hs.203961	ESTs, Moderately similar to AF116721 89	22.67	36.90
	438274	Al918906	Hs.55080	ESTs	1.00	1.00
	438378	AW970529	Hs.86434	hypothetical protein FLJ21816	38.92	38.00
	438403	AA806607	Hs.292206	ESTS	1.00 2.05	1.00 80.00
30	438494	AA908678	Hs.130183	ESTs	1.00	131.00
50	438546 438552	AW297204 AJ245820	Hs.125811 Hs.6314	type I transmembrane receptor (seizure-r	1.43	1.45
	438702	AI879064	Hs.54618	ESTs	1.00	34.00
	438724	AW612553	Hs.114670	Human DNA sequence from clone RP11-16L21	1.33	1.10
	438746	AI885815	Hs.184727	Human melanoma-associated antigen p97 (m	2.42	1.59
35	438779	NM_003787	Hs.6414	nucleolar protein 4	1.00	18.00
	438821	AA826425	Hs.192375	ESTs	2.03	2.57
	438885	A1886558	Hs.184987	ESTs	6.42	88.00
	438898	AA401369	Hs.190721	ESTs	22.41	17.00
40	438915	AA280174	Hs.285681	Williams-Beuren syndrome chromosome regi	1.00	1.00
40	438956	W00847	Hs.135056	Human DNA sequence from clone RP5-850E9	2.20 2.78	1.88 4.81
	439000 439023	AW979121 AA745978	Hs.28273	gb:EST391231 MAGE resequences, MAGP Homo ESTs	1.17	1.31
	439023	R96696	Hs.35598	ESTs	1.00	28.00
	439128	Al949371	Hs.153089	ESTs	1.00	67.00
45	439146	AW138909	Hs.156110	immunoglobulin kappa constant	1.38	1.41
	439223	AW238299	Hs.250618	UL16 binding protein 2	1.93	1.64
	439285	AL133916		hypothetical protein FLJ20093	46.23	139.00
	439318	AW837046	Hs.6527	G protein-coupled receptor 56	2.00	2.20
50	439343	AF086161	Hs.114611	hypothetical protein FLJ11808	6.10	7.37
50	439394	AA401369	Hs.190721	ESTs	3.39 1.83	17.00 3.07
	439410	AA632012	Hs.188746	ESTs heterochromatin-like protein 1	23.28	52.00
	439451 439452	AF086270 AA918317	Hs.278554 Hs.57987	B-cell CLL/lymphoma 11B (zinc finger pro	18.76	122.00
	439453	BE264974	Hs.6566	thyroid hormone receptor interactor 13	2.78	1.58
55	439477	W69813	Hs.58042	ESTs, Moderately similar to GFR3_HUMAN G	1.22	1.44
	439492	AF086310	Hs.103159	ESTs	7.46	39.00
	439523	W72348	Hs.185029	ESTs	1.00	1.19
	439592	AF086413	Hs.58399	ESTs	1.00	1.00
C 0	439606	W79123	Hs.58561	G protein-coupled receptor 87	33.61	1.00
60	439670	AF088076	Hs.59507	ESTs, Weakly similar to AC004858 3 U1 sm	1.00	1.00
	439702	AW085525	Hs.134182	ESTS	4.30 86.55	10.00 11.00
	439706	AW872527	Hs.59761 Hs.9598	ESTs, Weakly similar to DAP1_HUMAN DEATH sema domain, immunoglobulin domain (lg),	2.36	1.88
	439738 439750	BE246502 AL359053	Hs.57664	Homo sepiens mRNA full length Insert cDN	2.02	6.08
65	439759	AL359055	Hs.67709	Homo sapiens mRNA full length insert cDN	1.00	21.00
••	439780	AL109688	. 2.0	gb:Homo sapiens mRNA full length insert	7.27	25.00
	439840	AW449211	Hs.105445	GDNF family receptor alpha 1	1.00	1.00
	439926	AW014875	Hs.137007	ESTs	32.58	71.00
~~	439963	AW247529	Hs.6793	platelet-activating factor acetylhydrola	21.28	9.55
70	439979	AW600291	Hs.6823	hypothetical protein FLJ10430	68.83	61.00
	440006	AK000517	Hs.6844	hypothetical protein FLJ20510	1.83	4.02
	440028	AW473675	Hs.125843	ESTs, Weakly similar to T17227 hypotheti	1.42	2.54 54.00
	440106	AA864968	Hs.127699	KIAA1603 protein	1.00 24.18	54.00 52.00
75	440138 440273	AB033023 Al805392	Hs.318127 Hs.325335	hypothetical protein FLJ10201 Homo sapiens cDNA: FLJ23523 fis, clone L	3.21	4.72
, 5	440273	AW450991	Hs.192071	ESTs	38.63	113.00
	440325	NM_003812	Hs.7164	a disintegrin and metalloproteinase doma	62.88	147.00
	440492	R39127	Hs.21433	hypothetical protein DKFZp547J036	2.35	3.62
00	440527	AV657117	Hs.184164	ESTs, Moderately similar to S65657 alpha	10.84	57.00
80	440659	AF134160	Hs.7327	claudin 1	3.18	2.37
	440704	M69241	Hs.162	insulin-like growth factor binding prote	2.89	2.09
	440943	AW082298	Hs.146161	hypothetical protein MGC2408	2.02	1,41
	440994	A1160011	Hs.272068	ESTs	1.29 142.99	1.14 17.00
85	441020	AA401369 Al110684	Hs.190721	ESTs fibrinogen, B beta polypeplide	1.41	99.00
	441031	Al110684	Hs.7645	morningum a now portpopula	1171	

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	441128	AA570256	775	ESTs, Weakly similar to T23273 hypotheti	4.13	3.50
	441290	W27501	Hs.89605	cholinergic receptor, nicotinic, alpha p	1.00	1.00
	441362	BE614410	Hs.23044	RAD51 (S. cerevislae) homolog (E coli Re	130.23 22.03	43.00 1.00
5	441377 441390	BE218239 AI692560	Hs.202656 Hs.131175	ESTs ·	3.65	7.70
,	441497	R51064	Hs.23172	ESTs	1.00	1.00
	441525	AW241867	Hs.127728	ESTs	1.53	1.42
	441553	AA281219	Hs.121298	ESTs neuronal cell adhesion molecule	1.89 1.47	1,57 2,11
10	441607 441633	NM_005010 AW958544	Hs.7912 Hs.112242	normal mucosa of esophagus specific 1	216.22	363.00
10	441636	AA081846	Hs.7921	Homo sapiens mRNA; cDNA DKFZp566E183 (fr	2.31	2.05
	441737	X79449	Hs.7957	adenosine deaminase, RNA-specific	1.30	1.49
	441790	AA401369	Hs.190721	ESTs	44.15 1.00	17.00 1.00
15	441801 441919	AW242799 Al553802	Hs.86366 Hs.128121	ESTs ESTs	1.00	122.00
13	441937	R41782	Hs.22279	ESTs	0.86	1.37
	441954	AI744935	Hs.8047	Fanconi anemia, complementation group G	1.48	1.39
	442025	AW887434	Hs.11810	CDA11 protein	1.00 9.92	46.00 45.00
20	442029 442072	AW956698 AJ740832	Hs.14456 Hs.12311	neural precursor cell expressed, develop Homo sapiens clone 23570 mRNA sequence	25.05	77.00
20	442108	AW452649	Hs.166314	ESTs	3.61	3.14
•	442117	AW664964	Hs.128899	ESTs	3.00	5.49
	442137	AA977235	Hs.128830	ESTs, Weakly similar to Z192_HUMAN ZINC	1.00 1.92	1.00 1.66
25	442159 442179	AW163390 AA983842	Hs.278554 Hs.333555	heterochromatin-like protein 1 chromosome 2 open reading frame 2	27.22	50.00
23	442328	AI952430	Hs.150614	ESTs, Weakly similar to ALU4_HUMAN ALU S	5.00	3.42
	442432	BE093589	Hs.38178	hypothetical protein FLJ23468	181.59	76.00
	442530	A1580830	Hs.176508	Home sapiens cDNA FLJ14712 fis, clone NT	10.59 109.23	144.00 98.00
30	442547 442556	AA306997 AL137761	Hs.217484 Hs.8379	ESTs, Weakly similar to ALU1_HUMAN ALU S Homo sapiens mRNA; cDNA DKFZp586L2424 (f	1.00	53.00
50	442619	AA447492	Hs.20183	ESTs, Weakly similar to AF164793 1 prote	29.02	50.00
	442710	AI015631	Hs.23210	ESTs	1.00	19.00
	442717	R88362	Hs.180591	ESTs, Weakly similar to T23976 hypotheti	1.00 22.85	5.00 50.00
35	442875 442914	BE623003 AW188551	Hs.23625 Hs.99519	Homo sapiens clone TCCCTA00142 mRNA sequ hypothetical protein FLJ14007	25.33	82.00
55	442932	AA457211	Hs.8858	bromodomain adjacent to zinc finger doma	3.18	4.41
	442942	AW167087	Hs.131562	ESTs	8.45	64.00
	443068	Al188710	11- 00040	ESTS	1.00 1.00	27.00 24.00
40	443204 443211	AW205878 Al128388	Hs.29643 Hs.143655	Homo sapiens cDNA FLJ13103 fis, clone NT ESTs	12.42	2.00
70	443247	BE614387	Hs.333893	c-Myc target JPO1	128.84	96.00
	443324	R44013	Hs.164225	ESTs	0.02	4.59
	443383	A)792453	Hs.166507	ESTs	1.00 18.52	47.00 61.00
45	443400 443426	R28424 AF098158	Hs.250648 Hs.9329	ESTs chromosome 20 open reading frame 1	4.02	1.75
	443572	AA025610	Hs.9605	cleavage and polyadenylation specific fa	2.98	2.57
	443575	A1078022	Hs.269636	ESTs, Weakly similar to ALU1_HUMAN ALU S	1.00	29.00 16.00
	443614 443633	AV655386 AL031290	Hs.7645 Hs.9654	fibrinogen, B beta polypeptide similar to pregnancy-associated plasma p	1.00 1.00	39.00
50	443648	A1085377	Hs.143610	ESTs	39.81	70.00
	443715	AI583187	Hs.9700	cyclin E1	48.74	7.00
	443723	Al144442	Hs.157144	syntaxin 6	1.29 1.75	1.30 1.61
	443802 443859	AW504924 NM_013409	Hs.9805 Hs.9914	KIAA1291 protein foliistatin	1.35	1.13
55	443892	AA401369	Hs.190721	ESTs	1.00	17.00
- ·.	443947	W24187		gb:zb47f09.r1 Soares_fetal_lung_NbHL19W	1.33	1.64
	443991	NM_002250	Hs.10082	potassium intermediate/small conductance type I transmembrane protein Fn14	5.71 1.47	6.87 1.92
	444006 444009	BE395085 Al380792	Hs.10086 Hs.135104	ESTs	1.00	77.00
60	444017	U04840	Hs.214	neuro-oncological ventral antigen 1	1.00	1.00
	444127	N63620	Hs.13281	ESTs	1.00	29.00 1.00
200	444129 444279	AW294292 U62432	Hs.256212 Hs.89605	ESTs cholinergic receptor, nicotinic, alpha p	1.00 0.60	7.80
	444371	BE540274	Hs.239	forkhead box M1	2.91	1.14
65	444378	R41339	Hs.12569	ESTs	1.00	1.00
	444381	BE387335	Hs.283713	ESTs, Weakly similar to \$64054 hypotheti	469.00 12.88	556.00 105.00
	444461 444471	R53734 AB020684	Hs.25978 Hs.11217	ESTs, Weakly similar to 2109260A B cell KIAA0877 protein	24.91	90.00
	444489	AI151010	Hs.157774	ESTs	1.00	111.00
70	444619	BE538082	Hs.8172	ESTs, Moderately similar to A46010 X-lin	1.00	70.00
	444665	BE613126	Hs.47783	B aggressive tymphoma gene	30.56 1.00	139.00 1.00
	444707 444735	Al188613 BE019923	Hs.41690 Hs.243122	desmocollin 3 hypothetical protein FLJ13057 similar to	77.02	90.00
	444781	NM_014400	Hs.11950	GPI-anchored metastasis-associated prote	1.57	1.31
75	444783	AK001468	Hs.62180	anillin (Drosophila Scraps homolog), act	77.55	2.00
	445236	AK001676	Hs.12457	hypothetical protein FLJ 10814 ESTs	1.00 1.00	27.00 73.00
	445258 445413	A1635931 AA151342	Hs.147613 Hs.12677	CGI-147 protein	28.14	50.00
	445417	AK001058	Hs.12680	Homo sapiens cDNA FLJ10196 fis, clone HE	1.81	2.62
80	445443	AV653838	Hs.322971	ESTs	1.00	1.00
	445462	AA378776	Hs.288649	hypothetical protein MGC3077 hypothetical protein	2.0 9 1.87	1.70 70.00
	445517 445537	AF208855 AJ245671	Hs.12830 Hs.12844	EGF-like-domain, multiple 6	1.71	2.72
0.5	445580	AF167572	Hs.12912	skb1 (S. pombe) homolog	1.52	1.34
85	445654	X91247	Hs.13046	thioredoxin reductase 1	1.51	1.52

	w	O 02/086	443			
	445669	A1570830	Hs.174870	ESTs	10.95	11.45
	445818	BE045321	Hs.136017	ESTs	1.00	1.00
	445873	AA250970	Hs.251946	poly(A)-binding protein, cytoplasmic 1-	49.42 1.00	54.00 132.00
5	445885 445898	AI734009 AF070623	Hs.127699 Hs.13423	KIAA1603 protein Homo saplens clone 24468 mRNA sequence	1.00	1.00
	445903	Al347487	Hs.132781	class I cytokine receptor	1.00	36.00
	445932	BE046441	Hs.333555	Homo sapiens clone 24859 mRNA sequence	2.41	2.88
	445982	BE410233	Hs.13501	pescadillo (zebrafish) homolog 1, contai	1.60 1.00	1.35 42.00
10	446078 446102	Al339982 AW168067	Hs.156061 Hs.317694	ESTs ESTs	1.00	1.00
••	446157	BE270828	Hs.131740	Homo sapiens cDNA: FLJ22562 fis, clone H	1.70	1.53
	446269	AW263155	Hs.14559	hypothetical protein FLJ10540	73.01	48.00
	446292 446293	AF081497	Hs.279682 Hs.149722	Rh type C glycoprotein	1.55 1.00	1.26 2.00
15	446423	Al420213 AW139655	Hs.150120	ESTs ESTs	1.10	4.19
	446428	AW082270	Hs.12496	ESTs, Weakly similar to ALU4_HUMAN ALU S	0.53	3.26
	446432	Al377320	Hs.150058	ESTs	1.00	5.00
	446528 446574	AU076640 A)310135	Hs.15243 Hs.335933	nucleolar protein 1 (120kD) ESTs	1.36 3.89	1.31 72.00
20	446619	AU076643	Hs.313	secreted phosphoprotein 1 (osteopontin,	32.03	20.23
	446636	AC002563	Hs.15767	citron (rho-interacting, serine/threonin	4.19	5.07
	446783	AW138343	Hs.141867	ESTs	2.82 110.28	9.47 28.00
	446839 446849	BE091926 AU076617	Hs.16244 Hs.16251	mitotic spindle colled-coll related prot cleavage and polyadenylation specific fa	3.26	2.94
25	446856	AJ814373	Hs.164175	ESTs	6.38	11.30
	446872	X97058	Hs. 16362	pyrimidinergic receptor P2Y, G-protein c	1.98	2.03
	446880	AI811807	Hs.108646	Homo sapiens cDNA FLJ14934 fis, clone PL small inducible cytokine subfamily A (Cy	94.90 1.67	113.00 3.90
	446921 446989	AB012113 AK001898	Hs.16530 Hs.16740	hypothetical protein FLJ11036	2.82	3.12
30	447022	AW291223	Hs.157573	ESTs	1.00	170.00
	447033	Al357412	Hs.157601	ESTs	7.15 47.24	107.00 24.00
	447078 447081	AW885727 Y13896	Hs.9914 Hs.17287	ESTs polassium inwardly-rectifying channel, s	0.12	17.88
	447131	NM_004585	Hs.17466	retinoic acid receptor responder (tazaro	0.97	1.48
35	447149	BE299857	Hs.326	TAR (HIV) RNA-binding protein 2	1.24	1.26
	447153	AA805202	Hs.315562	ESTS	1.00 1.00	54.00 67.00
	447164 447178	AF026941 AW594641	Hs.17518 Hs.192417	Homo sapiens cig5 mRNA, partial sequence ESTs	3.42	50.00
	447250	A1878909	Hs.17883	protein phosphatase 1G (formerly 2C), ma	1.60	1.52
40	447289	AW247017	Hs.36978	melanoma antigen, family A, 3	1.00	1.00
	447342 447343	Al199268 AA256641	Hs.19322 Hs.236894	Homo sapiens, Similar to RIKEN cDNA 2010 ESTs, Highly similar to S02392 alpha-2-m	28.63 146.62	1.00 51.00
	447350	AJ375572	Hs.172634	ESTs	1.00	12.00
15	447377	N27687	Hs.334334	transcription factor AP-2 alpha (activat	2.55	63.00
45	447415 447425	AW937335 Al963747	Hs.28149 Hs.18573	ESTs, Weakly similar to KF3B_HUMAN KINES acylphosphatase 1, crythrocyte (common)	0.91 1.00	1.13 35.00
	447519	U46258	Hs.339665	ESTs	59.89	49.00
	447532	AK000614	Hs.18791	hypothetical protein FLJ20607	1.23	1.63
50	447534 447636	AA401369 Y10043	Hs.190721	ESTs high-mobility group (nonhistone chromoso	1,00 1,41	17.00 1.11
50	447688	N87079	Hs.19236	Target CAT	1.00	39.00
	447733	AF157482	Hs.19400	MAD2 (mitotic arrest deficient, yeast, h	1.17	1.12
	447769	AW873704	Hs.320831	Homo sapiens cDNA FLJ14597 fis, clone NT	6.47 0.73	5.95 2.34
55	447802 447850	AW593432 AB018298	Hs.161455 Hs.19822	ESTs SEC24 (S. cerevisiae) related gene famil	86.45	116.00
	447924	AI817226	Hs.313413	ESTs, Weakly similar to T23110 hypotheti	1.00	1.00
	447973	AB011169	Hs.20141	similar to S. cerevislae SSM4	3.50	4.27
	448030 448105	N30714 Al538613	Hs.325960 Hs.298241	membrane-spanning 4-domains, subfamily A Transmembrane protease, serine 3	4.13 1.15	142.00 2.24
60	448243	AW369771	Hs.52620	integrin, beta 8	15.84	1.00
	448278	WQ7369	Hs.11782	ESTs	0.97	1.90
	448290	AK002107	Hs.20843	Homo sapiens cDNA FLJ11245 fis, clone PL Homo sapiens cDNA FLJ14162 fis, clone NT	1.00 2.42	1.00 2.17
	448296 448357	BE622756 BE274396	Hs.10949 Hs.108923	RAB38, member RAS oncogene family	1.44	1.08
65	448390	AL035414	Hs.21068	hypothetical protein	1.00	43.00
	448469	AW504732	Hs.21275	hypothetical protein FLJ11011	2.63	2.49 2.53
	448569 448663	BE382657 BE614599	Hs.21486 Hs.106823	signal transducer and activator of trans hypothetical protein MGC14797	1.84 3.29	46.00
	448672	Al955511	Hs.225106	ESTs	1.00	21.00
70	448733	NM_005629	Hs.187958	solute carrier family 6 (neurotransmitte	1.82	1.08
	448741	BE614567	Hs.19574	hypothetical protein MGC5469 TATA box binding protein (TBP)-associate	2.48 23.53	1.92 20.00
	448757 448775	Al366784 AB025237	Hs.48820 Hs.388	nudix (nucleoside diphosphate linked moi	2.34	1.97
	448826	Al580252	Hs.293246	ESTs, Weakly similar to putative p150 [H	74.07	62.67
75	448830	AL031658	Hs.22181	hypothetical protein dJ310013.3	1.37	1.31 31.00
	448844 448988	Al581519 Y09763	Hs.177164 Hs.22785	ests gamma-aminobutyric acid (GABA) A recepto	1.00 1.84	1.95
	448993	AI471630	,	KIAA0144 gene product	1.63	1.49
90	449003	X76342	Hs.389	alcohol dehydrogenase 7 (class IV), mu o	1.00	1.00
80	449029 449040	N28989 AF040704	Hs.22891 Hs.149443	solute carrier family 7 (cationic amino putative turnor suppressor	1.97 0.97	2.26 1.56
	449048	Z45051	Hs.22920	similar to S68401 (cattle) glucose induc	27.13	90.00
	449053	A1625777	Hs.344766	EST6	8.33	44.00
85	449054	AF148848	Hs.22934	myoneurin G protein coupled receptor	73.85 2.58	104.00 27.00
02	449101	AA205847	Hs.23016	G protein-coupled receptor	2.00	21.00

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	449167	T05095	Hs.19597	KIAA1694 protein	1.61	2.36
	449207	AL044222	Hs.23255	nucleoporin 155kD	2.36	1.56
	· 449228	AJ403107	Hs.148590	protein related with psoriasts	1.15	1.15
5	449230	BE613348	Hs.211579	melanoma cell adhesion molecule	206.65	151.00
3	449305	A1638293	Un 70534	gb:tt09b07.x1 NCL_CGAP_GC6 Homo sapiens Homo sapiens, Similar to RIKEN cDNA 5730	17.28 26.39	45.00 35.00
	449318 449448	AW236021 D60730	Hs.78531 Hs.57471	ESTs	1.00	1.00
	449467	AW205006	Hs.197042	ESTs	1.00	1.00
	449523	NM_000579	Hs.54443	chemokine (C-C motif) receptor 5	56.80	216.86
10	449722	BE280074	Hs.23960	cyclin B1	150.03	1.00
	449976	H06350	Hs.135056	Human DNA sequence from clone RP5-850E9	2.16	2.85
	450001	NM_001044	Hs.406	solute carrier family 6 (neurotransmitte	1.17 1.79	1.45 2.38
	450098 450101	W27249 AV649989	Hs.8109 Hs.24385	hypothetical protein FLJ21080 Human hbc647 mRNA sequence	1.00	69.00
15	450149	AW969781	Hs.132863	Zic family member 2 (odd-paired Drosophi	1.00	1.00
	450193	AI916071	Hs.15607	Homo sapiens Fanconi anemia complementat	29.85	34.00
	450221	AA328102	Hs.24641	cytoskeleton associated protein 2	1.00	1.00
	450372	BE218107	Hs.202436	ESTs	1.00	1.00
20	450375	AA009647	Hs.8850	a disintegrin and metalloproteinase doma	51.26 123.20	93.00 181.00
20	450447 450568	AF212223 AL050078	Hs.25010 Hs.25159	hypothetical protein P15-2 Homo sapiens cDNA FLJ10784 fis, clone NT	1.00	19.00
	450589	AJ701505	Hs.202526	ESTs	1.00	23.00
	450684	AA872605	Hs.25333	interleukin 1 receptor, type II	1.00	100.00
~~	450701	H39960	Hs.288467	Homo sapiens cDNA FLJ12280 fis, clone MA	1.89	1.55
25	450705	U90304	Hs.25351	iroquois homeobox protein 2A (IRX-2A) (1.00	45.00
	450832	AA401369	Hs.190721	ESTs	25.17	17.00
	450937	R49131	Hs.26267 Hs.25740	ATP-dependant interferon response protei	90.92 3.33	90.00 1.70
	450983 451105	AA305384 Al761324	NS.23740	ERO1 (S. cerevisiae)-like gb:wi60b11.x1 NCI_CGAP_Co16 Homo sapiens	15.02	124.00
30	451110	A1955040	Hs.265398	ESTs, Weakly similar to transformation-r	1.00	143.00
	451253	H48299	Hs.26126	claudin 10	3.02	2.29
	451291	R39288	Hs.6702	ESTs	1.00	1.00
	451320	AW498974		dlacylglycerol kinase, zeta (104kD)	2.92	18.00
35	451380	H09280	Hs.13234	ESTS	6.90 35.75	6.67 72.00
33	451386 451437	AB029006 H24143	Hs.26334 Hs.31945	spastic paraplegia 4 (autosomal dominant hypothetical protein FLJ11071	1.00	69.00
	451462	AK000367	Hs.26434	hypothetical protein FLJ20360	1.83	2.10
	451524	AK001466	Hs.26516	hypothetical protein FLJ10604	1.13	1.07
40	451541	BE279383	Hs.26557	plakophilin 3	1.88	1.33
40	451592	Al805416	Hs.213897	ESTs	1.00	1.00
	451635	AA018899	Hs.127179	cryptic gene	1.52 4.95	1.92 17.00
	451743 451806	AA401369 NM_003729	Hs.190721 Hs.27076	ESTs RNA 3'-terminal phosphate cyclase	13.55	31.00
	451807	W52854	113.21010	hypothetical protein FLJ23293 similar to	1.55	35.00
45	451871	AI821005	Hs.118599	ESTs	1.81	2.53
	451952	AL120173	Hs.301663	ESTs	1.00	22.00
	452012	AA307703	Hs.279766	kinesin family member 4A	3.43	2.26
	452046	AB018345	Hs.27657	KIAA0802 protein	56.59 1.67	19.00 4.09
50	452194 452206	A!694413 AW340281	Hs.332649 Hs.33074	olfactory receptor, family 2, subfamily Homo sapiens, clone IMAGE:3606519, mRNA,	9.31	53.00
	452240	AA401369	Hs.190721		13.42	17.00
•	452256	AK000933	Hs.28661	Homo sapiens cDNA FLJ10071 fis, clone HE	39.03	94.00
	452281	T93500	Hs.28792	Homo sapiens cDNA FLJ11041 fis, clone PL	153.01	340.00
55	452291	AF015592	Hs.28853	CDC7 (cell division cycle 7, S. cerevisi	1.95 42.33	23.00
22	452295	BE379936	HS.28866	programmed cell death 10 ESTs, Weakly similar to S10590 cysteine	1.17	61.00 2.14
	452304 452340	AA025386 NM_002202	Hs.61311 Hs.505	ISL1 transcription factor, LIM/homeodoma	1.00	13.00
	452349	AB028944	Hs.29189	ATPase, Class VI, type 11A	1.09	1.42
	452367	U71207	Hs.29279	eyes absent (Drosophila) homolog 2	54.49	53.00
60 .	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro	1.00	32.00
	452410	AL133619	15-400400	Homo sapiens mRNA; cDNA DKFZp434E2321 (f	1.26	1.99
	452461 452571	N78223 W31518	Hs.108106 Hs.34665	transcription factor ESTs	24.47 54.61	35.00 102.00
	452613	AA461599	Hs.23459	ESTs	1.39	1.32
65	452699	AW295390	Hs.213062	ESTs	1.00	26.00
	452705	H49805	Hs.246005	ESTs	1.00	1.00
	452747	AF160477	Hs.61460	lg superfamily receptor LNIR	112.87	1.29
	452787	AW294022	Hs.222707	KIAA1718 protein	1.00	1.00
70	452795	AW392555	Hs.18878	hypothetical protein FLJ21620	1.00 7.91	1.00 75.00
70	452823 452833	AB012124 BE559681	Hs.30696 Hs.30736	transcription factor-like 5 (basic helix KIAA0124 protein	3.16	1.92
	452838	U65011	Hs.30743	preferentially expressed antigen in mela	174.35	1.00
	452862	AA401369	Hs.190721	ESTs	98.26	17.00
ac	452865	AW173720	Hs.345805	ESTs, Weakly similar to A47582 B-cell gr	1.55	1.00
75	452934	AA581322	Hs.4213	hypothetical protein MGC16207	1.73	1.19
	452946	X95425	Hs.31092	EphA5	1.00 1.58	1.00 1.98
	452976 453028	R44214 AB006532	Hs.101189 Hs.31442	ESTs RecQ protein-like 4	1.80	1.60
	453026 453095	AW295660	Hs.252756	ESTs	0.77	1.50
80	453102	NM_007197	Hs.31664	frizzled (Drosophila) homolog 10	1.00	1.00
	453103	Al301052	Hs.153444	ESTs	1.00	1.00
	453120	AA292891	Hs.31773	pregnancy-induced growth inhibitor	1.23	1.20
	453153	N53893	Hs.24360	ESTs	1.00 1.00	83.00 30.00
85	453160 453197	AJ263307 AJ916269	Hs.239884 Hs.109057	H2B histone family, member L ESTs, Weakly similar to ALU5_HUMAN ALU S	1.00	134.00
55	-W171	14010200	103001	Total transfer mine to transfer to the transfer		

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		AL133161	Hs.32360	hypothetical protein FLJ10867	1.69	1.93	101.0	.502/121/0
	453240	AI969564	Hs.166254	hypothetical protein DKFZp566i133	1.00	1.00		
	453317 453323	NM_002277 AF034102	Hs.41696 Hs.32951	keratin, hair, acidic,1 solute carrier family 29 (nucleoside tra	1.19 4.90	1.27 4.11		
5	453331	A1240665	Hs.8850	ESTs	199.42	340.00		
•	453392	U23752	Hs.32964	SRY (sex determining region Y)-box 11	1.00	16.00		
	453431	AF094754	Hs.32973	glycine receptor, beta	1.00	1.00		
	453439	AI572438	Hs.32976	guanine nucleotide binding protein 4	3.44 2.84	5.17 5.58		
10	453459 453563	BE047032 AW608906.co	Hs.257789	ESTs Hs.181163		protein MGC5629	4.58	90.00
10	453633	AA357001	Hs.34045	hypothetical protein FLJ20764	1.74	1.60		
	453775	NM_002916	Hs.35120	replication factor C (activator 1) 4 (37	19.49	1.00		
		AA534296	Hs.20953	ESTs	24.92	25.00		
15	453857 453867	AL080235 A1929383	Hs.35861 Hs.33032	DKFZP586E1621 protein hypothetical protein DKFZp434N185	167.59 1.00	66.00 39.00		
13	453883	A1638516	Hs.347524	cofactor required for Sp1 transcriptiona	1.97	1.58		
	453884	AA355925	Hs.36232	KIAA0186 gene product	63.89	20.00		
	453900	AW003582	Hs.226414	ESTs, Weakly similar to ALU8_HUMAN ALU S	20.41	16.00		
20	453922		Hs.36708	budding uninhibited by benzimidazoles 1	7.09 29.75	22.00 19.00		
20	453941 453964	U39817 Al961486	Hs.36820 Hs.12744	Bloom syndrome ESTs	1.00	1.00		
	453968	AA847843	Hs.62711	Homo saplens, clone IMAGE:3351295, mRNA	2.06	1.81		
	453976	BE463830	Hs.163714	ESTs	3.02	131.00		
25	454024	AA993527	Hs.293907	hypothetical protein FLJ23403	1.00	131.00		
23	454034 454042	NM_000691 T19228	Hs.575 Hs.172572	aldehyde dehydrogenase 3 family, member hypothetical protein FLJ20093	1.23 30.63	1.02 171.00		
	454059	NM_003154	Hs.37048	statherin	1.00	1.00		
	454066	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid	1.01	1.45		
20	454098	W27953	Hs.292911	ESTs, Highly similar to S60712 band-6-pr	1.26	1.11		
30	454241	BE144666	Un 110026	gb:CM2-HT0176-041099-017-c02 HT0176 Homo trinucleotide repeat containing 9	6.33 4.30	5.04 7.82		
	454417 454439	A1244459 AW819152	Hs.110826 Hs.154320	DKFZP56601646 protein	1.00	1.00		
		AW993247		gb:RC2-BN0033-180200-014-h09 BN0033 Homo	13.75	103.00	•	
25	455601	AI368680	Hs.816	SRY (sex determining region Y)-box 2	206.11	1.00		
35	456237	AA203682	11-02005	gb:zx52e07.r1 Soares_fetal_liver_spleen_	1.00	1.00		
	456321 456475	NM_001327 NM_000144	Hs.87225 Hs.95998	cancer/testis antigen Friedreich ataxia	1.14 1.00	1.10 48.00		
	456508	AA502764	Hs.123469	ESTs, Weakly similar to AF208855 1 BM-01	162.25	189.00		
40	456534	X91195	Hs.100623	phospholipase C, beta 3, neighbor pseudo	2.12	1.80		
40	456736	AW248217	Hs.1619	achaete-scute complex (Drosophila) homol	1.15	1.94		
	456759 456990	BE259150 NM_004504	Hs.127792 Hs.171545	delta (Drosophila)-like 3 HIV-1 Rev binding protein	1.00 16.42	1.00 84.00		
	457200	U33749	Hs.197764	thyroid transcription factor 1	0.57	1.76		
		AW968360	Hs.14355	Homo sapiens cDNA FLJ13207 fis, clone NT	2.71	4.15		•
45	457465	AW301344	Hs.122908	DNA replication factor	46.37	47.00		
	457489 457646	A1693815 AA725650	Hs.127179 Hs.112948	cryptic gene ESTs	1.12 1.55	1.35 2.51		
	457733		Hs.291971	ESTs	1.00	55.00		
~~		AA057484	Hs.35406	ESTs, Highly similar to unnamed protein	4.36	3.18		
50		BE545684	Hs.343566	KIAA0251 protein	1.00	1.32		
-	458098	BE550224 T28472	Hs.7655	metallothionein 1E (functional) U2 small nuclear ribonucleoprotein auxil	1.00 2.06	22.00 1.88		
	458207 458242	BE299588	Hs.28465	Homo sapiens cDNA: FLJ21869 fis, clone H	1.00	1.00		
	458247	R14439	Hs.209194	ESTs	7.00	9.85		
55		AW975460	Hs.142913	ESTs	1.00	3.00		
		AW451034	Hs.326525	arylsulfatase 0 RAN binding protein 1	1.31 1.98	2.01 1.71		•
	458933 459352	Al638429 AW810383	Hs.24763 Hs.206828	ESTs	12,60	63.00		. '
-a ·	459670		Hs.172004	tiin	1.00	1.00		
60	459702	A1204995	•	gb:an03c03.x1 Stratagene schizo brain S1	1.00	237.00		
	TABLE 98	3		•				
		-						
65	Pkey:			ntifier number		•		
		ber: Gene cluste		h			*	
•	Accession	n: Genoanka	ccession num	pers				
	Pkey	CAT Number	r Access	sion				
70	407746	10125_1	AK001	962 R69415 BE464605 AA418699 AA053293 AA1490	75 AA058396 AW33	8226 AW272659 AA45	4607 AJ 139535 A	W469852 Al275461
				982 AA730033 AA576507 AA991217 AA782067 AI98	5851 AA805864 AA!	505598 AW469857 R69	3546 AA988279 A	W001647 N63320
	408070	1036688_1		1 T27343 AA306950 AA360989 R58778 1852 BE350895				
	408660	107294_1		775 AA056342 Al538978 AW975281 AA664986				
75	409522	113735_1		382 AA075431				
	409866	1156522_1		2152 H41202 H29772				
	410032	1170435_1		985 BE065944 BE066008 BE066083 BE066093 454 AA713730 AA091294 AA584921 N86077 AW8367	781 AARN1N21 AAE7	QR76 AAEE1106 AAC2	3188 AWONEETT /	1955808 AI679388
	411089	123172_1		454 AA713730 AA091294 AA589921 NBBU77 AY8587 195 AA514764 AA454562 Al082382 AA595822 AA551				
80	411152	1234028_1	BF069	199 AW936012 AW877466 AW819782 AW935798 AW	/835546 AW936042	BE069121 AW835625	AW877536 AW93	35885 BE069202
		_	AW820	0019 AW935937 BE160180 AW935946 BE069101 BEI)69125 AW877527 E	3E160316 BE160398 A	W935794 AW835	701 AW935784
	412537	1304_1	AL031	778 X59711 NM_002505 M59079 A1870439 A1494259 3 Be079412 Be079428 N90322 A1631202 A1141758 A	AW664010 AA4050	63 AA436132 BE1745 JR62075 AI375230 AI2	10 AA412691 AI40 08445 AW235763	IUS 14 AA430UZ4 ALD44113 AA382556
			AW953	3 BEU/9412 BEU/9428 N30322 A651202 A1141730 A 3918 AA927051 AA889823 BE003094 AW390155 AW	360805 AW360823	W360810 AA425472	A1694282 ALD441	14 AI684577 AI809865

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	440044	400040 4	AI478773 AI160445 AI674630 N69088 AW665529 N49278 AI129239 AI457890 AI621264 AW297152 AI268215 AA907787 AI286170 AI017982 AI963541 AI469807 AI969353 BE552356 N66509 AA736741 AA382555 AW075811 AW292026
5	412811	132943_1	H06382 AW957730 AA352014 R13591 AA121201 D60420 BE263253 BE047862 Z41952 AI424991 AI693507 AI863108 AA599060 AI091148 AA598689 R39887 AA813482 AW016452 H06383 R41807 AI364268 AA620528 AI241940 AW089149 AW090733 AW088875 Z38240
3	413690	1383256_1	AA121202 R17734 BE157489 BE157560
	414883	15024_1	AA926960 AA926959 W76521 W24270 W21526 AA037172 BE267636 H83186 AA469909 N86396 AA001348 BE535736 AA081745 BE566245
			AA082436 H72525 H77575 N49786 W80565 H78746 BE569085 W04339 R98127 T55938 BE279271 AW960304 T29812 AA476873 BE297387
10			AA292753 AA177048 NM_001826 X54941 BE314366 AA908783 AI719075 BE270172 BE269819 AA889955 AI204630 W25243 AI935150 AA872039 W72395 T99630 AI422691 H98460 N31428 BE255916 H03265 AI857576 AA776920 AA910644 AA459522 AA293140 AW514667
10			R75953 AW662396 AA662522 Al865147 Al423153 AW262230 AA584410 AA583187 AW024595 AW069734 Al828996 AA282997 AA876046
			AW613002 AA527373 AW972459 AI831360 AA621337 AA100926 AA772418 AA594628 AI033892 W95096 AI034317 AA398727 AI085031
			N95210 AI459432 AI041437 AA932124 AA627684 AA935829 AI004827 AI423513 AI094597 H42079 R54703 AI630359 AA617681 AA978045
15			AA643280 W44561 AI991988 AI537692 AI090262 AA740817 AI312104 AI911822 AA416871 AI185409 AA129784 AA701623 AI075239 AI139549 AA633648 AI339996 AI336880 AA399239 AI078708 AI085351 AI362835 AI346618 AI146955 AI989380 AI348243 N92892 AA765850
			Al494230 Al278887 AA962596 Al492600 W80435 AA001979 R97424 Al129015 N24127 AA157451 AA235549 AA459292 AA037114 AA129785
			A1494211 AW059501 AW886710 R92790 N59755 AI361128 AW589407 H47725 H97534 H48076 H48450 T99631 AW300758 H03431 R76789
			AA954344 H77576 R96823 AI457100 N92845 N49682 H42038 BE220698 BE220715 H99552 AA701624 N74173 R54704 H79520 H72923 H03266 BE261919 AA769633 AA480310 AA507454 AA910586 AI203723 AW104725 W25611 W25071 T88980 H03513 T77589 R99156
20			W95095 R97470 AA702275 T77551 AA911952 H82956 N83673 AA283872
	415989	156454_1	AI267700 AI720344 AA191424 AI023543 AI469633 AA172056 AW958465 AA172236 AW953397 AA355086
	417324 418574	166714_1 17690_1	AW265494 AA455904 AA195677 AW265432 AW991605 AA456370 N28754 N28747 Al568146 Al979339 AA322671 AA322672 AW955043 Al990326 AA776406 Al016250 AA843678 AW451882 N23137 N23129
~~		1.000	W70051 AI038748 AA831327 AI925845 AW945895
25	418712	1784125_1	Z42183 T31621 T97478
	419443 419502	184788_1 18535_1	D62703 AA242966 D79798 AU076704 T74854 T74860 T72098 T73265 T73873 T69180 T74658 T58786 T60385 T73410 T68781 T67845 T67593 T73952 T67864 T60630
	410002	10005_1	T68367 T68401 T53959 T72360 T72099 T60377 T58961 T71712 T72821 T64738 T74645 T72037 T68688 T72063 T73258 T72826 T64242
20			T68220 T74673 T71800 T68355 T61227 T62738 T69317 T53850 T64692 T73768 T73962 T73382 T68914 T70975 T73400 T60631 T73277
30			T73203 T70498 T61409 T58925 NM_000508 M64982 T68301 T73729 T69445 T60424 T67922 T67736 T68716 T67755 T74765 T73819 T58719 T74756 T60477 T74863 T61109 T68329 T58850 T71857 T73425 T53736 T68607 T58898 T64309 T72031 T72079 T64305 T71908 T68107
			T71916 T73787 T56035 T64425 T71870 T60476 T61376 T67820 T71895 T41006 T69441 T68170 T74617 T71958 T69440 T61875 R06796
			H48353 T71914 T53939 T64121 AA693996 T72525 T67779 T68078 AA011465 AA345378 AV654847 AV654272 AV656001 Al064740 T82897
35			N33594 AA344542 AW805054 Al207457 T61743 AA026737 H94389 AA382695 AA918409 T68044 S82092 T39959 Al017721 AA312395 AA312919 T40156 H66239 AV652989 H38728 R98521 AV655200 R95790 W03250 W00913 AA344136 AV660126 R97923 AA343596
-			AW470774 AV651256 N54417 AA812862 AW182929 Al111192 H61463 H72060 AA344503 H38639 Al277511 AV661108 Al207625 T47810
			AA235252 T27853 T47778 R95746 H70620 AA701463 AW827166 R98475 C20925 AV657287 T71959 T71313 T73920 T73333 T61618 T69293
			T69283 T73931 T72178 T72456 AV645639 AV653476 T72957 T72300 T58906 T71457 T70494 T72956 T70495 T68267 T74407 T85778 AA344726 T27854 T74485 T74101 T73868 T71518 T72304 AA343853 T73909 T68070 T72065 H72149 T73493 T73495 AV645993 R02293
40			T70475 T64751 AA344441 AA343657 AA345732 AA344328 Al110639 AA344603 AF063513 T64696 T68516 T72223 T60507 T67633 R29500
	•		T72517 R02292 T60599 T69206 T70452 T74577 R29366 T61277 T74914 T60352 R29675 T74843 AV645792 AA344408 T69197 T72057
			T69368 T69358 T68258 AV650429 T73341 T61702 T74598 T40095 K02272 T40106 AA343045 AA341908 AA341907 AA342807 AA341964 T53747 T72042 T62764 A\064899 AA343060 T67832 T72440 T71770 T68091 T69108 T72449 T69167 T71289 T68251 A\0654844 T64375
4.0		•	AA345234 T67598 AA011414 T68036 H48262 Al207557 T68219 W86031 T69081 T64232 R93196 T62136 AV650539 H67459 T72978
45			AA344583 T60362 H58121 T95711 T72803 T68055 T71715 R29036 T72793 T69122 T64595 T62888 T69139 T68291 T64652 T67971 T46862
			AA693592 Al248502 R29454 T64764 T57001 T73052 T71429 T51176 T58866 AV655414 H90426 AA342489 T73666 T67848 T72512 T53835 T67837 T73317 T74273 T69420 T68245 T74380 T67862 T74474 T56068
	419936	189181_1	Al792788 BE142230 AA252019
50 `	421582	2041_1	AI910275 X00474 X52003 X05030 NM_003225 AA314326 AA308400 AA506787 AA314825 AI571948 AA507595 AA614579 AA587613 R83818 AA568312 AA614409 AA307578 AI925552 AW950155 AI910083 M12075 BE074052 AW004668 AA578674 AA582084 BE074053 BE074126
50	••.	*	BE074140 AA514776 AA588034 BE074051 BE074068 AW009769 AW050690 AA858276 R55389 AI001051 AW050700 AW750216 AA614539
	2400400		BE074045 Al307407 AW602303 BE073575 Al202532 AA524242 Al970839 Al909751 BE076078 Al909749 R55292
	422128 423034	211994_1 224122_1	AW881145 AA490718 M85637 AA304575 T06067 AA331991 AL119930 AA320696 AW752565
55	423816	23234_1	AL031985 AL137241 Al792386 Al733664 Al857654 Al049911
	424200	236595_1	AA337221 AA336756 AW966196
	424999 · 426966	245835_1 273896_1	AW953120 R56325 AA349562 AI493134 AI498691 AW771508 AI498457 AI768408 AI783624 AI383985 AI580267 D79813 AA393768
	426991	27415_1	AK001536 AA191092 AW510354 AI554256 AL353968 AA134266
60	427260	276598_1	AA663848 AA400100 AA401424
	428023	28589_2	AL038843 AA161338 BE268213 AA425597 N87306 AA092969 BE566038 AA247451 N47392 AI928802 AW182584 AW027872 AI819831 AI936994 W56258 AI653448 AI278611 AI283557 AI824306 AW338658 AW150899 AA687514 N47393 N29885 AA973469 AI038904 AI292064
			AI034339 AW674593 N72156 AI079733 AI038683 AI291616 AA491599 AA993675 AA837380 BE006554 BE006473 AI087090 T33044
65			AA652043 AI203503 AA583959 W35283 AI129926 Z41844 AW020925 AW575848 AI684603 AA493297 AI140689 AI277175 AA425444
.03	429220	301384_1	AI932767 W02632 BE396786 R37261 AW207206 AW341473 AA448195 AI951341
	429978	31150_1	AA249027 AL038984 AK001993 AL080066 AV652725 BE566226 AA345557 AA315222 AA090585 AA375688 AA301092 AA298454 W05762
			AW607939 H51658 D83880 N84323 BE296821 AW947007 D61461 AW079261 AA329482 AW901780 AI354442 AA772275 R31663 AI354441
70			AI767525 H92431 AI916735 H93575 AI394255 AW014741 AI573090 C06195 AW612857 AW265195 AI339558 AI377532 AI308821 AI919424 AI589705 AW055215 AI336532 AI338051 AA806547 C76509 C00618 AW071172 AW769904 AA630381 AI678018 AI863985 D79662 BE221049
			AW265018 AI589700 AW196655 N76573 AI370908 BÈ042393 N75017 AI698870 AW960115
	430439 430935	31808_1 325772_1	AL133561 AL041090 AL117481 AL122069 AW439292 AI968826 AW072916 AI184913 AA489195 AW466994 AW469044 N59350 AI819642 AI280239 AI220572 AA789302 AI473611 AW841126 D60937
	430935 431089	327825_1	AWU72916 A1164913 AA469193 AW460994 AW460944 NO9330 A1619642 A1260239 A1220572 AA789302 A1473611 AW841126 D00937 BE041395 AA491826 AA621946 AA715980 AA666102
75	431322	331543_1	AW970622 AA503009 AA502998 AA502989 AA502805 T92188
	432407	34624_1	AA221036 R87170 BE537068 BE544757 C18935 AW812058 T92565 AA227415 AA233942 AA223237 AA668403 AA601627 AW869639 REGS1823 BEG00570 AM861470 AW847510 AA388A2 AW824832 AW848698 COA690 AA387244 AW824637 AA068730
			BE051833 BE000620 AW961170 AW847519 AA308542 AW821833 AW945688 C04699 AA205504 AA377241 AW821667 AA055720 AW817981 AW856468 AA155719 AA179928 T03007 AW754298 AA227407 AA113928 AA307904 C16859
οΛ	434414	38585_1	AI798376 S46400 AW811617 AW811616 W00557 BE142245 AW858232 AW861851 AW858362 AA232351 AA218567 AA055556 AW858231
80			AW857541 AW814172 H66214 AW814398 AF134164 AA243093 AA173345 AA199942 AA223384 AA227092 AA227080 T12379 AA092174 T61139 AA140775 AA600929 AA19774 AA197756 AA197756 AA19776 AA19776 AA19776 AA19776 AA19776 AA19776 AA19776
			T61139 AA149776 AA699829 AW879188 AW813567 AW813538 Al267168 AA157718 AA157719 AA100472 AA100774 AA130756 AA157705 AA157730 AA157715 AA053524 AW849581 AW84566 C05254 AW882836 T92637 AW812621 AA206583 AA209204 BE156909 AA226824
			AI829309 AW991957 N66951 AA527374 H66215 AA045564 AI694265 H60808 AA149726 AW195620 BE081333 BE073424 AW817662
85	436608	42361_3	AW817705 AW817703 AW817659 BE081531 H59570 AA628980 A1126603 BE504035
-	70000	4500 I_U	r v vacanda na nadada Madalada

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	438091	44964_1		AW373052 T55662 Al299190 BE174210 AW579001 H01811 W40186 R67100 Al923886 AW952164 AA628440 AW898607 AW898616 AA709126 AW898528 AW898544 AA947932 AW898625 AW898622 Al276125 Al185720 AW510698 AA987230 T52522 BE467708 AW243400 AWM33642 Al788745 Al186932 D52654 D55017 D52715 D52477 D53933 D54679 Al298739 Al146984 Al922204 N98343 BE174213 AA845571
5			•	A1813854 A1214518 A1635262 A1139455 A1707807 A1698085 AW884528 A1024768 A1004723 AW087420 A1565133 N94964 A1268939 AW513280 A1061126 A1435818 A1859106 A1360506 A1024767 AA513019 AA757598 X56196 AA902959 A1334784 A1860794 AA010207 AW890091 AW513771 A1951391 A1337671 T52499 AA890205 A1640908 H75966 AA463487 AA358688 A1961767 A1866295 AA780994 A1985913 BE174196 AA029094 AW592159 T55581 N79072 A1611201 AA910812 A1220713 AW149306 A1758412 AA045713 R79750 N76096
10	439000 439285	467716_1 47065_1		AW979121 AA847986 AA829098 AL133916 N79113 AF086101 N76721 AW950828 AA364013 AW955684 AI346341 AI867454 N54784 AI655270 AI421279 AW014882 AA775552 N62351 N59253 AA626243 AI341407 BE175639 AA456968 AI358918 AA457077
	439780 441128	47673_1 51021_2		AL 109688 R23665 R26578 AA570256 AW014761 AA573721 AI473237 AI022165 AA554071 AA127551 N90525 AW973623 AA447991 AA243852 BE328850 AI148171 AI359627 AI005068 AI356567 AA232991 AW016855 AA906902 AA233101 AA127550 BE512923
15	443068	558874_1		A1188710 A1032142 AW078833 N30308 AW675632 A1219028 A1341201 N22181 H95390
	443947 447636	586160_1 7301_1	,	W24187 W24194 R17789 Y10043 NM_005342 L05085 AL034450 BE614226 AW749053 AA379173 AA248230 BE514634 AA334622 R70656 AA367593 AA214649 AA369318 AW957081 R05760 AA039903 AI886597 AW630122 AA906264 AA041527 R01145 AI088688 BE463637 AA398795 AI354883 AI768938 AI569996 AI452952 AI168582 AI189869 AI086670 AW262560 AW613854 AA862839 AA435840 AA670197 AI024032 AI990659 AI990089 N81095 AA847919 AW960150 AA211075 AA044704 AA367594 AW582587 AW858854 AW818630 AW818281 AW818433 AW582595
20	448993	79225_1		AA096002 N83992 AI471630 BE540637 BE265481 AW407710 BE513882 BE546739 AA053597 BE140503 BE218514 AW956702 AI656234 AI636283 AI567265 AW340858 BE207794 AA053085 R69173 AA292343 AA454908 AA293504 AI659741 AI927478 AA399460 AI760441 AA346416 BE047245 AA730380 AA394063 AA454833 AI982791 AI567270 AI813332 AI767858 AA427705 D20284 AI221458 BE048537 AI263048 AA346417
25	449305	804424_1		AA911497 BE537702 Al638293 AW813561
٠	451105 451320	859083_1 86576_1		AI761324 AW880941 AW880937 AW118072 AI631982 T15734 AA224195 AI701458 W20198 F26326 AA890570 N90552 AW071907 AI671352 AI375892 T03517 R88265 AI124088 AA224388 AI084316 AI354686 T33652 AI140719 AI720211 T03490 AI372637 T15415 AW205836 AA630384 T03515 T33230
30	451807 .	8865_1	1	AA017131 AA443303 T33623 Al222556 T33511 T33785 Al419606 D55612 W52854 AL117600 BE208116 BE208432 BE206239 BE082291 AW953423 AA351619 BE180648 BE140560 W60080 AA865478 N90291 AW450652 AW449519 AA993634 Al806539 AA351618 AW449522 Al827626 AA904788 AA380381 AA886045 AA774409 BE003229 Z41756
	452410	9163_1		ANASOS ANASOS ANASOS IS ACESSOS AIGUSTOS ANASOS ANAS
35				AI269603 AI565082 AI807095 AI476629 AA505909 AI368449 AI686077 AI582930 AW085038 AA757863 AA730154 AI767072 AA468316
	454241 455175	1067807_1 1257335_1	1	A1734130 A1734138 AA426284 AA433997 A1741241 AW043563 A1732741 A1732734 AA437369 AA425820 AA664048 R74130 BE144666 BE184942 AW238414 BE184946 AW993247 AW861464
40	456237 458098	168730_1 47395_1	ĺ	AA203682 R11958 BE550224 AA832519 N45402 AW885857 N29245 BE465409 W07677 AW970089 A1299731 AA482971 BE503548 H18151 W79223 AF086393 AA461301 W74510 R34182 A1090689 N46003 BE071550 R28075 AW134982 A124024 A1138906 AW026179 A1572316 BE466182 A1205395
				A!276154 A!273269 A!422817 A!371014 A!421274 A!188525 AA939164 BE549810 AW137865 A!694996 BE503841 AA459718 BE327407 BE467534 BE218421 BE467767 AA989054 BE467063 A!797130 BE327781
45				
	TABLE 9C			
	Pkey: Ref:	Sequence	SOUTCE.	orresponding to an Eos probeset . The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA
50	Strand: Nt_position:	sequence Indicates	of huma DNA stra	an chromosome 22.° Dunham I. et al., Nature (1999) 402:489-495. and from which exons were predicted. de positions of predicted exons.
	Pkey	Ref	Strand	Nt_position
55	400512 400517	9796593 9796686	Minus Minus	1439-1615 49996-50346
	400560 400664	9843598 8118496	Plus Plus	94182-94323,97056-97243,101095-101236,102824-103005 13558-13721,13942-14090,14554-14679
60	400665 400666	8118496 8118496	Plus Plus	16879-17023 17982-18115,20297-20456
00	400749	7331445	Minus	9162-9293
	400763 401027	8131616 7230983	Minus Minus	35537-35784 70407-70554,71060-71160
65	401093 401203	8516137 9743387	Minus Minus	22335-23166 172961-173056,173868-173928
	401212 401411	9858408 7799787	Plus Minus	87839-88028 144144-144329
	401435	8217934	Minus	54508-55233
70	401464 401714	6682291 6715702	Minus Plus	170688-170834 96484-96681
	401747 401760	9789672 9929699	Minus Plus	118596-118816,119119-119244,119609-119761,120422-120990,130161-130381,130468-130593,131097-131258,131866- 131932,132451-132575,133580-134011 83126-83250,85320-85540,94719-95287
75	401780	7249190	Minus	28397-28617,28920-29045,29135-29296,29411-29567,29705-29787,30224-30573 83215-83435,83531-83656,83740-83901,84237-84393,84955-85037,66290-86814
13	401781 401785	7249190 7249190	Minus Minus	165776-165996,166189-166314,166408-166569,167112-167268,167387-167469,168634-168942
	401797 401961	6730720 4581193	Plus Minus	6973-7118 124054-124209
80	401985 401994	2580474 4153858	Plus Minus	61542-61750 42904-43124,43211-43336,44607-44763,45199-45281,46337-46732
50	402075	8117407	Plus	121907-122035,122804-122921,124019-124161,124455-124610,125672-126076
	402260 402265	3399665 3287673	Minus Plus	113765-113910,115653-115765,116808-116940 21059-21168
85	402297 402408	6598824 9796239	Plus Minus	35279-35405,35573-35659 110326-110491

	W	O 02/08	6443	
	402420	9796339	Plus	129750-129919
	402674	8077108	Minus	39290-39502
	402802	3287156	Minus	53242-53432
	402994	2996643	Minus	4727-4969
5	403137	9211494	Minus	92349-92572,92958-93084,93579-93712,93949-94072,94591-94748,95214-95337
_	403306	8099945	Plus	127100-127251
	403329	8516120	Plus	96450-96598
	403381	9438267	Minus	26009-26178
	403478	9958258	Plus	116458-116564
10	403485	9966528	Plus	2888-3001,3198-3532,3655-4117
- •	403627	8569879	Minus	23868-24342
	403715	7239669	Plus	85128-85292
	404044	9558573	Minus	225757-225939
	404076	9931752	Minus	3848-3967
15	404101	8076925	Minus	125742-125997
	404140	9843520	Plus	37761-38147
	404165	9926489	Minus	69025-69128
	404185	4572584	Minus	129171-129327
	404210	5006246	Plus	169926-170121
20	404253	9367202	Minus	55675-56055
	404287	2326514	Plus	53134-53281
	404298	9944263	Minus	73591-73723
	404347	9838195	Plus	74493-74829
	404440	7528051	Plus	80430-81581
25	404721	9856648	Minus	173763-174294
	404794	4826439	Plus	101619-101898
	404854	7143420	Plus	14260-14537
	404877	1519284	Plus	1095-2107
	404927	7342002	Plus	68690-69563
30	404996	6007890	Plus	37999-38145,38652-38998,39727-39872,40557-40674,42351-42450
	405449	7622497	Plus	42236-42570
	405568	6006906	Plus	35912-36065
	405572	3800891	Plus	85230-85938
	405646	4914350	Plus	741-969
35	405676	4557087	Plus	73195-73917
	405770	2735037	Plus	61057-62075
	405932	7767812	Minus	123525-123713
	406137	9166422	Minus	30487-31058
	406360	9256107	Minus	7513-7673
40	406399	9256288	Minus	63448-63554
-	406467	9795551	Plus	182212-182958
				•

- TABLE 10A: Potential Therapeutic, Diagnostic and Prognostic targets for Therapy of Lung Cancer and Non-malignant Lung Disease
 Table 2A shows about 307 genes up-regulated in non-malignant lung disease relative to lung tumors and normal body tissues and/or down-regulated in lung tumors relative to normal lung and non-malignant lung disease. These genes were selected from about 59680 probesets on the Eos/Affyrnetrix Hu03 Genechip array.
- Table 108 show the accession numbers for those Pkey's lacking UnigenelD's for table 10A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.
- Table 10C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 10A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Pkey: 1 Unique Eos probeset identifier number ExAccn: Exemplar Accession number, Genbank accession number

Unigene Title: Unigene number Unigene gene title

R1: Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the

average of normal lung samples

R2: Average of non-malignant lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples

65	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
	404394		. 7	ENSP00000241075:TRRAP PROTEIN.	0.79	3.10
	404916			Tarpet Exon	1.00	159.00
	405257			Target Exon	1.00	422.00
	407228	M25079	Hs.155376	hemoglobin, beta	0.47	2.33
70	407568	AA740964	Hs.62699	ESTs	1.00	123.00
	408562	AI436323	Hs.31141	Homo sapiens mRNA for KIAA1568 protein,	1.00	230.00
	409031	AA376836	Hs.76728	ESTs	1.00	128.00
	410434	AF051152	Hs.63668	toll-like receptor 2	39.65	149.00
	410467	AF102546	Hs.63931	dachshund (Drosophila) homolog	1.00	109.00
75	410808	T40326	Hs.167793	ESTs	1.14	13.14
	412351	AL135960	Hs.73828	T-cell acute lymphocytic leukemia 1	0.37	2.27
	412372	R65998	Hs.285243	hypothetical protein FLJ22029	1.00	173.00
	413795	AL040178	Hs.142003	ESTs	0.10	11.90
	414154	AW205314	Hs.323060	ESTs	0.62	2.09
80	414214	D49958	Hs.75819	glycoprotein M6A	0.03	4.55
	414998	NM_002543	Hs.77729	oxidised low density lipoprotein (lectin	0.64	2.97
	415122	D60708	Hs.22245	ESTs	0.07	8.97
	415765	NM 005424	Hs.78824	tyrosine kinase with immunoglobulin and	0.67	1.65
	415775	H00747	Hs.29792	ESTs, Weakly similar to 138022 hypotheti	0.29	2.64
85	415910	U20350	Hs.78913	chemokine (C-X3-C) receptor 1	1.00	145.00

	W	O 02/086	443			
	416319	AI815601	Hs.79197	CD83 antigen (activated 8 lymphocytes, i	15.32	237.00
	416402	NM_000715	Hs.1012	complement component 4-binding protein,	0.64	4.00
	417355	D13168	Hs.82002	endothelin receptor type B	0.01	3.90
5	417421	AL138201	Hs.82120	nuclear receptor subfamily 4, group A, m	36.30	357.00
)	417511 418489	AL049176 U76421	Hs.82223 Hs.85302	chordin-like adenosine deaminase, RNA-specific, B1 (h	1.00 0.02	179.00 6.00
	418726	BE241812	Hs.87860	protein tyrosine phosphatase, non-recept	1.00	113.00
	418741	H83265	Hs.8881	ESTs, Wealdy similar to S41044 chromosom	0.44	1.90
10	418883	BE387036	Hs.1211	acid phosphatase 5, tartrate resistant	0.96	2.04
10	419086	NM_000216	Hs.89591	Kallmann syndrome 1 sequence	0.62 0.03	2.74 6.90
	419150 419235	T29618 AW470411	Hs.89640 Hs.288433	TEK tyrosine kinase, endothelial (venous neurotrimin	1.48	5.13
	419407	AW410377	Hs.41502	hypothetical protein FLJ21276	37.55	336.00
	420556	AA278300	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L	0.80	3.65
15	420656	AA279098	Hs.187636	ESTs	1.65	8.07
	420729	AW964897	Hs.290825	ESTs Homo sapiens mRNA; cDNA DKFZp586N0121 (f	2.99 0.46	25.82 1.95
	421177 422060	AW070211 R20893	Hs.102415 Hs.325823	ESTs, Moderately similar to ALU5_HUMAN A	1.00	156.00
	422426	W79117	Hs.58559	ESTs	0.03	7.44
20	422652	AW967969	Hs.118958	syntaxin 11	0.14	3.62
	423099	NM_002837	Hs.123641	protein tyrosine phosphatase, receptor t	0.01 0.75	3.16 141.75
	424433 424585	H04607 AA464840	Hs.9218 Hs.131987	ESTs ESTs	1.00	167.00
	424711	NM_005795	Hs.152175	calcitonin receptor-like	0.43	3.01
25	424973	X92521	Hs.154057	matrix metalloproteinase 19	0.37	19.45
	425023	AW956889	Hs.154210	endothelial differentiation, sphingolipi	0.14	3.35
	425664	AJ006276	Hs.159003	transient receptor potential channel 6	1.00 0.68	94.00 1.42
	425998 426657	AU076629 NM_015865	Hs.165950 Hs.171731	fibroblast growth factor receptor 4 solute carrier family 14 (urea transport	0.03	3.74
30	426753	T89832	Hs.170278	ESTs	1.00	141.00
•	427558	D49493	Hs.2171	growth differentiation factor 10	1.00	117.00
	427983	M17706	Hs.2233	colony stimulating factor 3 (granulocyte	0.75	2.20 2.25
	428467 428927	AK002121 AA441837	Hs.184465 Hs.90250	hypothetical protein FLJ11259 ESTs	0.76 0.01	3.62
35	429496	AA453800	Hs.192793	ESTs	1.00	138.00
	430468	NM_004673	Hs.241519	anglopoietin-like 1	1.00	132.00
	431385	BE178536	Hs.11090	membrane-spanning 4-domains, subfamily A	1.00	157.00
	431728	NM_007351	Hs.268107	multimerin ESTs, Highly similar to AF175283 1 zinc	1.00 0.34	157.00 2.24
40	431848 432128	Al378857 AA127221	Hs.126758 Hs.117037	ESTs	0.00	1.15
. •	432519	Al221311	Hs.130704	ESTs, Weakly similar to BCHUIA S-100 pro	0.01	2.06
	433043	W57554	Hs.125019	lymphoid nuclear protein (LAF-4) mRNA	1.00	267.00
	433803	AI823593	Hs.27688	ESTs	1.00 1.05	105.00 3.15
45	434730 435472	AA644669 AW972330	Hs.193042 Hs.283022	ESTs triggering receptor expressed on myelold	0.83	1.94
15	436532	AA721522	, 13.200022	gb:nv54h12.r1 NCI_CGAP_Ew1 Homo sapiens	1.00	218.00
	437119	Al379921	Hs.177043	ESTs	1.00	133.00
	437140	AA312799	Hs.283689	activator of CREM in testis	0.67 1.00	122.67 142.00
50	437211 437960	AA382207 AI669586	Hs.5509 Hs.222194	ecotropic viral integration site 2B ESTs	1.00	147.00
50	438202	AW169287	Hs.22588	ESTs	1.00	141.00
	438873	Al302471	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L	0.71	3.66
	438875	AA827640	Hs.189059	ESTs	23.32	370.00
55	441048 441188	AA913488 AW292830	Hs.192102 Hs.255609	ESTs ESTs	0.77 3.43	8.50 16.36
55	441499	AW298235	Hs.101689	ESTs	1.00	167.00
	444513	AL120214	Hs.7117	glutamate receptor, lonotropic, AMPA 1	1.00	151.00
	444527	NM_005408	Hs.11383	small inducible cytokine subfamily A (Cy	46.47	153.00
60	444561 445279	NM_004469 R41900	Hs.11392 Hs.22245	c-fos induced growth factor (vascular en ESTs	0.01 0.60	3.08 141.00
00	446017	N98238	Hs.55185	ESTs	0.18	2.39
	446984	AB020722	Hs.16714	Rho guanine exchange factor (GEF) 15	0.10	2.16
	446998	N99013	Hs.16762	Homo sapiens mRNA; cDNA DKFZp564B2062 (f	0.01	2.53
65	447357	A1375922 A1800470	Hs.159367 Hs.171941	ESTs ESTs	0.46 18.05	2.64 296.00
05	448106 448253	H25899	Hs.201591	ESTS	1.00	141.00
•	449275	AW450848	Hs.205457	periaxin	0.56	1.38
	450400	AI694722	Hs.279744	ESTs	0.88	4.33
70	450696	A1654223	Hs.16026	hypothetical protein FLJ23191	0.52 0.79	2.08 2.01
70	450726 451497	AW204600 H83294	Hs.250505 Hs.284122	retinolo acid receptor, alpha Wnt inhibitory factor-1	0.35	2.03
	451533	NM_004657	Hs.26530	serum deprivation response (phosphatidyl	0.13	2.25
	453636	R67837	Hs.169872	ESTs	1.00	116.00
75	458332	AI000341	Hs.220491	ESTS	1.00 0.20	192.00 2.98
13	459580 400269	AA022888	Hs.176065	ESTs Eos Control	0.40	2.40
	403421			NM_016369*:Homo sapiens claudin 18 (CLDN	0.53	1.77
	407570	Z19002	Hs.37096	zinc finger protein 145 (Kruppel-like, e	0.01	3.18
80	412295	AW088826	Hs.117176	poly(A)-binding protein, nuclear 1	0.56	1.74
OU.	414517 417204	M24461 N81037	Hs.76305 Hs.1074	surfactant, pulmonary-associated protein surfactant, pulmonary-associated protein	0.64 0.33	1.50 1.16
	418307	U70867	Hs.83974	solute carrier family 21 (prostaglandin	0.53	1.55
	418935	T28499	Hs.89485	carbonic anhydrase IV	0.20	1.28
85	421502	AF111856	Hs.105039	solute carrier family 34 (sodium phospha	0.78	1.90
O)	421798	N74880	Hs.29877	N-acylsphingosine amidohydrolase (acid c	0.59	1.54

	**	/ ^ 02/086	143			
	423354	O 02/086/ AB011130	Hs.127436	calcium channel, voltage-dependent, alph	0.59	1.55
	423738	AB002134	Hs.132195	airway trypsin-like protease	10.14	51.00
	425211	M18667	Hs.1867	progastricsin (pepsinogen C)	0.35	1.62
_	425438	T62216	Hs.270840	ESTs	0.23	9.45
5	426828	NM_000020	Hs.172670	activin A receptor type II-like 1	0.03	1.71 1.49
	427019 428043	AA001732 T92248	Hs.173233 Hs.2240	hypothetical protein FLJ10970 uteroglobin	0.01 0.42	1.26
	430280	AA361258	Hs.237868	Interleukin 7 receptor	0.46	2.43
	431433	X65018	Hs.253495	surfactant, pulmonary-associated protein	0.57	1.59
10	431723	AW058350	Hs.16762	Homo saplens mRNA; cDNA DXFZp564B2062 (f	0.29	1.80
	432985	T92363	Hs.178703	ESTs	0.32	2.27
	441835	AB036432	Hs.184	advanced glycosylation end product-speci	0.31 0.55	1.51 1.78
	442275 443709	AW449467 A1082692	Hs.54795 Hs.134662	ESTS ESTS	0.00	3.02
15	444325	AW152618	Hs.16757	ESTs	0.32	2.49
	450954	AJ904740	Hs.25691	receptor (calcitonin) activity modifying	0.46	1.74
	451558	NM_001089	Hs.26630	ATP-binding cassette, sub-family A (ABC1	0.52	1.87
	453310	X70697	Hs.553	solute carrier family 6 (neurotransmilte	0.00	3.30
20	456855	AF035528	. Hs.153863	MAD (mothers against decapentaplegic, Dr	0.01 0.66	2.31 2.20
20	444342 400754	NM_014398	Hs.10887	similar to lysosome-associated membrane Target Exon	1.00	297.00
	401045			C11001883*:gi[6753278]ref[NP_033938.1] c	1.00	109.00
	401083			NM_016582*:Homo sapiens peptide transpor	0.89	1.39
0.5	402474			NM_004079:Homo saplens cathepsin S (CTSS	1.45	4.47
25	402808			ENSP00000235229:SEMB.	1.00	1.87
	403021			C21000030:gi 9955960 ref NP_063957.1 AT	1.00 1.06	149.00 2.96
	403438 403687			NM_031419*:Homo sapiens molecule possess NM_007037*:Homo sapiens a disintegrin-li	0.04	4.89
	403764			NM_005463:Homo sapiens heterogeneous nuc	1.00	225.00
30	404277			NM_019111*:Homo sapiens major histocompa	0.97	1.93
	404288			NM_002944*:Homo sapiens v-ros avian UR2	1.00	68.00
	404518	Al815601		CD83 antigen (activated B lymphocytes, i	0.02	1.83
	405106			C11001637*:gi 5032241 ref NP_005732.1 z	1.00 1.00	235.00 93.00
35	405381 406387			Target Exon Target Exon	1.37	6.02
55	406646	M33600		major histocompatibility complex, class	0.86	2.46
	406714	Al219304	Hs.266959	hemoglobin, gamma G	0.01	3.19
	406753	AA505665	Hs.217493	annexin A2	1.00	147.00
40	406973	M34996	Hs.198253	major histocompatibility complex, class	1.03	2.04
40	407248	U82275	Hs.94498	teukocyte immunoglobulin-like receptor,	1.00 1.00	64.00 90.00
	407510 407731	U96191 NM_000066	Hs.38069	gb:Human trophoblast hypoxia-regulated f complement component 8, beta polypeptide	1.00	67.00
	407830	NM_001086	Hs.587	arylacetamide deacetylase (esterase)	1.00	102,00
	408045	AW138959	Hs.245123	ESTs	1.00	70.00
45	408074	R20723		ESTs	1.00	112.00
	408374	AW025430	Hs.155591	forkhead box F1	0.07	10.17
	409064 409083	AA062954 AF050083	Hs.141883	ESTs interleukin 12A (natural killer cell sti	0.39 1.00	2.31 95.00
	409063	W03754	Hs.673 Hs.50813	hypothetical protein FLJ20022	0.01	4.55
50	409203	AA780473	Hs.687	cytochrome P450, subfamily IVB, polypept	0.01	3.72
	409238	AL049990	Hs.51515	Homo sapiens mRNA; cDNA DKFZp564G112 (fr	1.00	79.00
	409389	AB007979	Hs.301281	Homo sapiens mRNA, chromosome 1 specific	0.14	27.35
	409718	D86640	Hs.56045	src homology three (SH3) and cysteine ri	1.00 0.64	113,00 2.47
55	410798 411020	BE178622 NM_006770	Hs.16291 Hs.67726	gb:PM3-HT0605-270200-001-a02 HT0605 Homo macrophage receptor with collagenous str	0.55	2.40
55	41 1667	BE160198	113.07720	gb:QV1-HT0413-010200-059-h03 HT0413 Homo	1.00	111.00
	412000	AW576555	Hs.15780	ATP-binding cassette, sub-family A (ABC1	1.00	95.00
	412358	BE047490	Hs.24172	ESTs	1.00	87.00
60	412420	AL035668	Hs.73853	bone morphogenetic protein 2	1.43	8.07 3.07
60	412564	X83703	Hs.31432	cardiac ankyrin repeat protein CXC chemokine ligand 16	0.02 0.93	3.07 1.72
	412869 412870	AA290712 N22788	Hs.82407 Hs.82407	CXC chemokine ligand 16	0.97	1.51
	413529	U11874	Hs.846	interleukin 8 receptor, beta	0.02	2.42
	413533	BE146973		gb:QV4-HT0222-011199-019-e05 HT0222 Homo	0.65	1.50
65 °	413689	BE157286	Hs.20631	zinc finger protein, subfamily 1A, 5 (Pe	20.87	232.00
	413724	AA131466	Hs.23767	hypothetical protein FLJ 12666	1.00	80.00
	413800	AI129238	Hs.192235	ESTs ESTs, Weakly similar to S65657 alpha-1C-	1.00 1.00	85.00 213.00
	413802 413829	AW964490 NM_001872	Hs.32241 Hs.75572	carboxypeptidase B2 (plasma)	0.02	3.93
70	414376	BE393856	Hs.66915	ESTs, Weakly similar to 16.7Kd protein (1.00	115.00
	414577	AI056548	Hs.72116	hypothetical protein FLJ20992 similar to	0.49	1.94
	414700	H63202	Hs.38163	ESTs	0.03	3.75
	415078	AA311223	Hs.283091	found in inflammatory zone 3	0.86	1.95
75	415120	N64464 BE260352	Hs.34950	ESTs neutrophil cytosolic factor 2 (65kD, chr	1.00 0.60	120.00 2.48
, ,	415323 415335	BE269352 AA847758	Hs.949 Hs.111030	ESTs	1.00	95.00
	415582	W92445	Hs. 165195	Homo sapiens cDNA FLJ14237 fis, clone NT	1.00	136.00
	416030	H15261	Hs.21948	ESTs	0.02	8.07
90	416427	BE244050	Hs.79307	Rac/Cdc42 guanine exchange factor (GEF)	1.00	73.00
80	416464	NM_000132	Hs.79345	coagulation factor VIII, procoagulant co	0.70	3.36
	416585	X54162 L43821	Hs.79386 Hs.80261	leiomodin 1 (smooth muscle) enhancer of filamentation 1 (cas-like do	0.06 0.70	6.56 3.66
	416847 417148	AA359896	Hs.80261 Hs.293885	hypothetical protein FLJ14902	1.00	114.00
	417370	T28651	Hs.82030	tryptophanyl-IRNA synthetase	0.85	1.30
85	417673	T87281	Hs.16355	ESTs	0.15	15.54

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	418067	Al127958	Hs.83393	cystatin E/M	0.81	1.74
	418296	C01566	Hs.86671	ESTa	1.00	99.00
	418643	J03798	Hs.86948	small nuclear ribonucleoprotein D1 polyp	1.00	60.00
5	418832	X04011	Hs.88974	cytochrome b-245, beta polypeptide (chro	2.40 0.67	14.74 · 3.16
5	418945 419261	BE246762 X07876	Hs.89499 Hs.89791	arachidonate 5-lipoxygenase wingless-type MMTV integration site fami	1.00	73.00
	419564	U08989	Hs.91139	solute carrier family 1 (neuronal/epline	1.00	192.00
	419574	AK001989	Hs.91165	hypothetical protein	1.00	94.00
10	419968	X04430	Hs.93913	interleukin 6 (interferon, beta 2)	61.16	500.00
10	420256	U84722 AA258124	Hs.76206 Hs.293878	cadherin 5, type 2, VE-cadherin (vascula ESTs, Moderately similar to ZN91_HUMAN Z	0.52 1.00	1.70 172.00
	420285 420577	AA278436	Hs.186649	ESTs	1.00	97.00
•	421262	AA286746	Hs.9343	Homo sapiens cDNA FLJ14265 fis, clone PL	1.00	64.00
1.5	421445	AA913059	Hs.104433	Homo sapiens, clone iMAGE:4054868, mRNA	0.88	1.51
15	421470	R27496	Hs.1378	annexin A3	0.05 1.00	11.26 73.00
	421478 421563	AI683243 NM_006433	Hs.97258 Hs.105806	ESTs, Moderately similar to S29539 ribos granulysin	0.82	2.42
	421566	NM_000399	Hs.1395	early growth response 2 (Krox-20 (Drosop	5.50	31.57
	421855	F06504	Hs.27384	ESTs, Moderately similar to ALU4_HUMAN A	1.00	129.00
20	421913	A1934365	Hs.109439	osteoglycin (osteoinductive factor, mime	1.00	101.00
	421952	AA300900	Hs.98849	ESTs, Moderately similar to AF161511 1 H	0.60 1.00	63.60 148.00
	422232 422386	D43945 AF105374	Hs.113274 Hs.115830	transcription factor EC heparan sulfate (glucosamine) 3-O-sulfot	1.40	3.98
	423168	R34385	Hs.124940	GTP-binding protein	0.34	3.59
25	423196	AK001866	Hs.125139	hypothetical protein FLJ11004	0.55	2.00
	423387	AJ012074		vasoactive intestinal peptide receptor 1	0.09	2.13
	423424	AF150241	Hs.128433	prostaglandin D2 synthase, hematopoietic	1.00 1.00	141.00 66.00
	423456 423696	AL110151 Z92546	Hs.128797	DKFZP586D0824 protein Sushi domain (SCR repeat) containing	0.73	1.27
30	424027	AW337575	Hs.201591	ESTs	0.54	2.58
-	424212	NM_005814	Hs.143131	almostolain A22 (transmembrane)	0.77	2.47
	425087	R62424	Hs.126059		1.00	74.00
	425175	AF020202	Hs.155001	UNC13 (C. elegans)-like	0.85 1.18	1.96 2.56
35	425771 426486	BE561776 BE178285	Hs.159494 Hs.170056	Bruton agammaglobulinemia tyrosine kinas Homo sapiens mRNA; cDNA DXFZp586B0220 (f	1.00	76.00
33	427507	AF240467	Hs.179152	toll-like receptor 7	1.00	63.00
	427618	NM_000760	Hs.2175	colony stimulating factor 3 receptor (gr	0.60	2.19
	427732	NM_002980	Hs.2199	secretin receptor	0.97	1.42
40	427952	AA765368	Hs.293941	ESTs, Moderately similar to A53959 throm	1.00	105.00 80.00
40	428709	BE268717	Hs.104916	hypothetical protein FLJ21940 ESTs	1.00 0.09	2.55
	428769 428780	AW207175 Al478578	Hs.106771 Hs.50636	ESTs	1.00	98.00
	428833	AI928355	Hs.185805	ESTs	1.00	113.00
	429657	D13626	Hs.2465	KIAA0001 gene product; putative G-protei	1.00	52.00
45	430212	AA469153		gb:nc67f04.s1 NCI_CGAP_Pr1 Homo sapiens	1.00	132.00
	430226	BE245562	Hs.2551	adrenergic, beta-2-, receptor, surface	0.11 1.00	15.60 103.00
	430376 430414	AW292053 AW365665	Hs.12532 Hs.120388	chromosome 1 open reading frame 21 ESTs	0.50	6.96
	430656	AA482900	Hs.162080	ESTs	1.00	70.00
50	430843	Al734149	Hs.119514	ESTs	1.00	90.00
	430998	AF128847	Hs.204038	indolelhylamine N-methyltransferase	0.29	1.84
	431217	NM_013427	Hs.250830	Rho GTPase activating protein 6 ESTs	1.00 0.91	79.00 1.67
	431921 432176	N46466 AW090386	Hs.58879 Hs.112278	arrestin, beta 1	0.66	2.63
55	432203	AA305746	Hs.49	macrophage scavenger receptor 1	1.00	76.00
	432231	AA339977	Hs.274127	CLST 11240 protein	0.46	1.46
	432485	N90866	Hs,276770	CDW52 antigen (CAMPATH-1 antigen)	0.79	2.25
	432522	D11466	Hs.51	phosphatidylinositol glycan, class A (pa matrilin 3	1.93 0.04	4.83 5.79
60	432596 432850	AJ224741 X87723	Hs.278461 Hs.3110	angiolensin receptor 2	1.00	167.00
•	433138	AB029496	Hs.59729	semaphorin sem2	0.04	9.16
	433563	Al732637	Hs.277901	ESTS	1.00	91.00
	433588	Al056872	Hs.133386	ESTs	120.16	315.00
65	434445	A1349306	Hs.11782	ESTs ESTs, Weakly similar to transformation-r	- 0.60 1.00	1.84 128.00
05	435496 435974	AW840171 U29690	Hs.265398 Hs.37744	Homo sapiens bela-1 adrenergic receptor	1.00	108.00
	436061	A1248584	Hs.190745	Homo sapiens cDNA: FLJ21326 fis, clone C	1.00	91.00
	437157	BE048860	Hs.120655	ESTs	1.00	87.00
70	437207	T27503	Hs.15929	hypothetical protein FLJ12910	1.00	105.00
70	437311	AA370041	Hs.9456	SWI/SNF related, matrix associated, acti	1.00 1.00	71.00 115.00
	437439 438199	H29796 AW016531	Hs.269622 Hs.122147	ESTs ESTs	1.00	80.00
	439551	W72062	Hs.11112	ESTs	0.30	3.10
	440515	AJ131245	Hs.7239	SEC24 (S. cerevisiae) related gene famil	1.00	77.00
.75	440887	AI799488	Hs.135905	ESTs	1.00	85.00
	441025	AA913880	Hs.176379	ESTs	1.00 0.79	82.00 1.89
	441384 441735	AA447849 AI738675	Hs.288660 Hs.127346	Homo sapiens cDNA: FLJ22182 fis, clone H ESTs	1.00	75.00
	442200	AW590572	Hs.235768	ESTs	0.78	5.83
80	442832	AW206560	Hs.253569	ESTs	0.03	10.88
	442957	Al949952	Hs.49397	ESTs	1.00	70.00
	443282	T47764	Hs.132917	ESTs	1.00 1.00	197.00 253.00
	443547 443951	AW271273 F13272	Hs.23767 Hs.111334	hypothetical protein FLJ12666 ferritin, light polypeptide	0.55	2.09
85	444330	AI597655	Hs.49265	ESTs	1.00	90.00
2-						

Plus

7636-8054

6006920

85

405381

TABLE 11A: Genes Distinguishing Adenocardinoma from Other Lung Diseases and Normal Lung

Table 11A shows about 84 genes upregulated in lung adenocarcinomas relative to other lung tumors, non-malignant lung disease, and normal lung. These genes were selected from about 59680 probesets on the Eos/Affyrnetrix Hu03 Genechlp array.

Table 11B show the accession numbers for those Pkey's lacking UnigenelD's for table 11A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarly using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 11C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in lable 11A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Pkey:

Unique Eos probeset Identifier number Exemplar Accession number, Genbank accession number Unigene number 15 ExAcon:

UnigenelD: Unigene Title: R1:

5

10

Unigene gene title

Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the

average of normal lung samples

Average of normally and lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples 20 R2:

20	R2:	Averag	e of non-mailg	ant lung disease samples (including bronchitis, emphysema, tibrosis, atelectasis,				
	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2		
	403329		••	Target Exon	1.00	61.00		
_	406399			NM_003122*:Homo sapiens serine protease	1.00	39.00		
25	406690	M29540	Hs.220529	carcinoembryonic antigen-related cell ad	226.37	350.00		
	407869	AI827976	Hs.24391	hypothetical protein FLJ13612	0.77	1.18		
	407881	AW072003	Hs.40968	heparan sulfate (glucosamine) 3-O-sulfot	1.00	10.00		
	408908	BE296227	Hs.250822	serine/threonine kinase 15	7.76	1.00		
<u>.</u>	409103	AF251237	Hs.112208	XAGE-1 protein	80.44	40.00		
30	409187	AF154830	Hs.50966	carbamoyl-phosphale synthetase 1, mitoch	1.00	1.00		
	409269	AA576953	Hs.22972	hypothetical protein FLJ13352	1.00	1.00		
	410076	T05387	Hs.7991	ESTs	1.12	1.50		
	410102	AW248508	Hs.279727	Homo sapiens cDNA FLJ14035 fis, clone HE	9.89	1.00		
25	410399	BE068889		synuclein, gamma (breast cancer-specific	0.92	1.06		
35	411908	L27943	Hs.72924	cytidine deaminase	1.00 1.02	1.00 1.03		
	412612	NM_000047	Hs.74131	arylsulfatase E (chondrodysplasia puncta	0.84	1.03		
	414075	U11862	Hs.75741	amilioride binding protein 1 (amine oxida ESTs, Weakly similar to MUC2_HUMAN MUCIN	3.67	1.00		
	416208 417542	- AW291168 J04129	Hs.41295 Hs.82269	progestagen-associated endometrial prote	1.28	1.35		
40	419183	U60669	Hs.89663	cytochrome P450, subfamily XXIV (vitamin	1.00	1.00		
70	419502	AU076704	113.03000	fibrinogen, A alpha polypeptide	13.05	115.00		
	419631	AW188117	Hs.303154	popeye protein 3	1.00	13.00		
	420931	AF044197	Hs.100431	small inducible cytokine B subfamily (Cy	1.00	8.00		
	421155	H87879	Hs.102267	lysyl oxidase	1.00	15.00		
45	421190	U95031	Hs.102482	mucin 5, subtype B, tracheobronchial	1.17	1.55		
-	421474	U76362	Hs.104637	solute carrier family 1 (glutamate trans	1.46	1.76		
	421515	Y11339	Hs.105352	GalNAc alpha-2, 6-sialyltransferase I, I	1.00	3.00		
	421582	Al910275		trefoil factor 1 (breast cancer, estroge	1.23	1.00		
	422026	U80736	Hs.110826	trinucleotide repeat containing 9	1.00	52.00		
50	422095	AI868872	Hs.282804	hypothetical protein FLJ22704	4.37	2.34		
	422311	AF073515	Hs.114948	cytokine receptor-like factor 1	1.15	1.78		
	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	1.69	3.17		
	423472	AF041260	Hs.129057	breast carcinoma amplified sequence 1	48.13	72.00		
55	423554	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	1.00	50.00		
23	424502	AF242388	Hs.149585	lengsin	1.00 1.00	1.00 59.00		
	424544	M88700	Hs.150403 Hs.153704	dopa decarboxylase (aromatic L-amino ad	21.35	1.00		
	424905 424960	NM_002497 BE245380	Hs.153704 Hs.153952	NIMA (never in mitosis gene a)-related k 5' nucleotidase (CD73)	1.00	1.00		
	425523	AB007948	Hs.158244	KIAA0479 protein	1.00	35.00		
60	426230	AA367019	Hs.241395	protease, serine, 1 (trypsin 1)	1.00	83.00		
O,O	427701	AA411101	Hs.243886	nuclear autoantigenic sperm protein (his	7.41	34.00		
	428585	AB007863	Hs.185140	KIAA0403 protein	1.00	6.00		
	428758	AA433988	Hs.98502	hypothetical protein FLJ14303	1.06	1.13		
	429170	NM_001394	Hs.2359	dual specificity phosphatase 4	16.18	105.00		
65	429263	AA019004	Hs.198396	ATP-binding cassette, sub-family A (ABC1	1.07	1.00		
	429610	AB024937	Hs.211092	LUNX protein; PLUNC (palate lung and nas	1.59	1.69		
	430508	AI015435	Hs.104637	ESTs	4.75	7.27		
	430985	AA490232	Hs.27323	ESTs, Weakly similar to 178885 serine/th	0.94	1.28		
70	431548	AI834273	Hs.9711	novel protein	5.66	15.00		
70	431566	AF176012	Hs.260720	J domain containing protein 1	49.76	37.00		
	431986	AA536130	Hs.149018	Novel human gene mapping to chomosome 20	1.19	1.47 1.06		
	432375	BE536069	Hs.2962	S100 calcium-binding protein P	1.65 1.00	48.00		
	432677	NM_004482	Hs.278611	UDP-N-acetyl-alpha-D-galactosamine:polyp	1.00	19.00		
75	433556	W56321	Hs.111460	calcium/calmodulin-dependent protein kin	3.71	8.00		
15	433819	AW511097 AW950905	Hs.112765 Hs.3697	ESTs serine (or cysteine) proteinase inhibito	29.31	72.00		
	434001 434424	AVV930903 AI811202	Hs.325335	Homo sapiens cDNA: FLJ23523 fis, clone L	1.00	64.00		
	434792	AA649253	Hs.132458	ESTs	8.52	44.00		
	436217	T53925	Hs.107	fibrinogen-like 1	57.97	31.00		
80	436749	AA584890	Hs.5302	lectin, galactoside-binding, soluble, 4	1.10	1.41		
-	436972	AA284679	Hs.25640	claudin 3	1.59	1.46		
	437866	AA156781		metallothioneln 1E (functional)	3.62	101.00		
	437935	AW939591	Hs.5940	mucin 13, epithelial transmembrane	1.60	1.39		
0.5	438915	AA280174	Hs.285681	Williams-Beuren syndrome chromosome regi	1.00	1.00		
85	439451	AF086270	Hs.278554	heterochromatin-like protein 1	23.28	52.00		

				•			
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	439759 441031	AL359055 Al110684	Hs.67709 Hs.7645	Homo sapiens mRNA full length insert cDN fibrinogen, B beta polypeptide	1.00 1.41	21.00 99.00	
	441377	BE218239	Hs.202656	ESTs	22.03	1.00	
-	443614	AV655386	Hs.7645	fibrinogen, B beta polypeptide	1.00	16.00	
5	443813 443991	AA876372 NM_002250	Hs.93961 Hs.10082	Homo sapiens mRNA; cDNA DKFZp667D095 (fr potassium intermediate/small conductance	1.20 5.71	1.99 6.87	
	444670	H58373	Hs.332938	hypothetical protein MGC5370	1.98	38.00	
	444931	AV652066	Hs.75113	general transcription factor IIIA	1.00	54.00	
10	446102 446163	AW168067 AA026880	Hs.317694 Hs.25252	ESTs Homo sapiens cDNA FLJ13603 fis, clone PL	1.00 1.00	1.00 36.00	
10	446469	BE094848	Hs.15113	homogentisate 1,2-dioxygenase (homogenti	1.00	11.00	
	447388	AW630534	Hs.76277	Homo sapiens, clone MGC:9381, mRNA, comp	1.24	1.16	
	447532 448243	AK000614 AW369771	Hs.18791 Hs.52620	hypothetical protein FLJ20607 integrin, beta 8	1.23 15.84	1.63 1.00	
15	448844	Al581519	Hs.177164	ESTs	1.00	31.00	
	449444	AW818436	Hs.23590	solute carrier family 16 (monocarboxylic	1.00	83.00	
	451807 452689	W52854 F33868	Hs.284176	hypothetical protein FLJ23293 similar to transferrin	1.55 1.54	35.00 1.44	
	4523392	U23752	Hs.32964	SRY (sex determining region Y)-box 11	. 1.00	16.00	
20	453464	AI884911	Hs.32989	receptor (catcitonin) activity modifying	1.55	2.45	5.
	453735	A1066629	Hs.125073	ESTs	1.01	1.30	
	TABLE 1	1B					
25	Disease	I belove Co	bt-!do	ntifier number			•
23	Pkey: CAT num	ber: Gene clust		numer number			
	Accession		ccession num	bers	•		•
	Dless	CAT Number	r Access	lon			
30	Pkey 410399	11995_1	BE068	889 BE068882 AF044311 AF017256 NM_003087 A	F037207 AF010	126 AA633976 AA	872836 BE298825 BE299889 Al016464 Al684600
			Al9365	27 AA804675 AA394097 AI139933 AA946606 BE17	'1313 AA72240	7 AA293803 A1468	480 AA056035 AA055968 AW796957 Al637713
	440502	10525 4.	AA410	737 H49348 AA486472 AA411094 AA235594 AA407 704 T74854 T74860 T72098 T73265 T73873 T6918	2624 AA443638 N T74658 T687	1 AW452137 AA421 RE TENSRE T73410	708 AW255211 AI493256 AA365132 AW966044 TER781 TE7845 TE7593 T73952 TE7864 TE0630
	419502	18535_1·	T68367	704 174834 174680 172033 173263 173673 10316 7 T68401 T53959 T72360 T72099 T60377 T58961 T	71712 T72821	T64738 T74645 T7	2037 T68688 T72063 T73258 T72826 T64242
35			T68220) T74673 T71800 T68355 T61227 T62738 T69317 T	53850 T64692	T73768 T73962 T7	3382 T68914 T70975 T73400 T60631 T73277
				T70498 T61409 T58925 NM_000508 M64982 T683 T60477 T74863 T61109 T68329 T58850 T71857 T			
				5 T73787 T56035 T64425 T71870 T60476 T61376 T			
40			H48353	3 T71914 T53939 T64121 AA693996 T72525 T6777	9 T68078 AA01	1465 AA345378 A\	/654847 AV654272 AV656001 Al064740 T82897
40			N33594	1 AA344542 AW805054 A1207457 T61743 AA02673 919 T40156 H66239 AV652989 H38728 R98521 AV	7 H94389 AA38	32695 AA918409 TI	58044 S82092 T39959 AI017721 AA312395
			AA312 AW470	919 140156 M66239 AV652969 N36726 K96321 AV 1774 AV651256 N54417 AA812862 AW182929 A111	1192 H61463 H	72060 AA344503 F	H38639 Al277511 AV661108 Al207625 T47810
			AA2352	252 T27853 T47778 R95746 H70620 AA701463 AW	827166 R9847	5 C20925 AV65728	7 T71959 T71313 T73920 T73333 T61618 T69293
45			T69283	T73931 T72178 T72456 AV645639 AV653476 T72	957 T72300 T5	8906 T71457 T704	94 T72956 T70495 T68267 T74407 T85778
43	•		AA344	726 T27854 T74485 T74101 T73868 T71518 T7230 5 T64751 AA344441 AA343657 AA345732 AA34432	8 Al110639 AA:	3909 168070 1720 344603 AF063513	T64696 T68516 T72223 T60507 T67633 R29500
			T72517	R02292 T60599 T69206 T70452 T74677 R29366 1	61277 T74914	T60352 R29675 T7	74843 AV645792 AA344408 T69197 T72057
			T69368	T69358 T68258 AV650429 T73341 T61702 T7459	3 T40095 K022	72 T40106 AA3430	45 AA341908 AA341907 AA342807 AA341964
50			153747 44345	7 T72042 T62764 Al064899 AA343060 T67832 T724 234 T67598 AA011414 T68036 H48262 Al207557 T0	4U 17777U 108 SR219 WREN31	1091 169108 17244 T69081 T64232 R9	9 169167 17 1269 166251 AV654644 164575 13196 T62136 AV650539 H67459 T72978
50			AA344	583 T60362 H58121 T95711 T72803 T68055 T7171	5 R29036 T727	93 T69122 T64595	T62888 T69139 T68291 T64652 T67971 T46862
			AA693	592 Al248502 R29454 T64764 T57001 T73052 T714	129 T51176 T58	1866 AV655414 H9	0426 AA342489 T73666 T67848 T72512 T53835
	421582	2041_1	T67837	'	74474 156068 AA308400 AA5		A1571048 AA507505 AA614579 AA587613 R83818
55	421302	2041_1	AA568	79 X00474 X52003 X03030 NIM_003223 AAS14320 312 AA614409 AA307578 AI925552 AW950155 AI9	10083 M12075	BE074052 AW0046	668 AA578674 AA582084 BE074053 BE074126
	,		BE074	140 AA514776 AA588034 BE074051 BE074068 AW	009769 AW050	690 AA858276 R5	5389 A1001051 AW050700 AW750216 AA614539
4.	. 407000	44422.0	BE0740	045 A1307407 AW602303 BE073575 A1202532 AA5: 781 AW293839 U52054 AA024963 AA778446 BE07	24242 A1970839	9 A1909751 BE0760)78 A 909749 R55292 34040 BE164012 BE163072 BE163074 BE163902
ير من	437866	44433_2	AA 136	781 AW293839 US2US4 AAU24983 AA776446 BEU7 481 AW468444 BE185091 AW468002 AA687333 AA	NB11830 AA581	806 Al866686 Al57	2124 AA043777 AA040926 D20160 AI536733
60			AA8124	189 AW874142 AI4718B3 W84421 AA156850		•	
	451807	8865_1	W5285	4 AL117600 BE208116 BE208432 BE206239 BE08: 1652 AW449519 AA993634 AI806539 AA351618 AW	2291 AW95342 1449522 A18276	3 AA351619 BE18(38 AA904788 AA?	ID48 BE140560 W60080 AA8654/8 N90291 80381 AAR86045 AA774400 RE003229 741756
			AVV45U	010 DECAM ECCOUNT PCOCEENA E1 CEPPTIA NU	443044 MIO410	IEU PASONI GO PANS	0000 1410000TO 1411 1700 OLUMEED ET 1100
65	74B1 = -	40 :					
UJ	TABLE 1	10					,
	Pkey:	Unique nur	mber correspo	nding to an Eos probeset			F - 1 11 3.17 - 11 11/1 1 - 11/1 PASA
	Ref:			digit numbers in this column are Genbank Identifier mosome 22." Dunham I. et al., Nature (1999) 402:4		"Dunham I. et al." r	eters to the publication entitled "The UNA
70	Strand:			mosome 22. Dunnam I. et al., Nature (1999) 402.44 m which exons were predicted.	13- 1 3J.		
	Nt_position			tions of predicted exons.			
	•		-				

Pkey 403329 406399

75

Ref 8516120 9256288 Strand Plus Minus Nt_position 96450-96598 63448-63554

PCT/US02/12476 WO 02/086443

TABLE 12A: Genes Distinguishing Squamous Cell Cardinoma from Other Lung Diseases and Normal Lung

Table 12A shows about 72 genes upregulated in squamous cell carcinomas of the lung relative to other lung turnors, non-mailgnant lung disease, and normal lung. These genes were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array.

Table 12B show the accession numbers for those Pkey's lacking UnigenelD's for table 12A. For each probaset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Ctustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 12C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 12A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

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Pkey: Unique Eos probeset identifier number ExAcon:

Exemplar Accession number, Genbank accession number

UnigenelD:

Unigene number

Unigene Title: 20

Unigene gene title
Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the R1:

average of normal lung samples

ples (including bronchilis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples DΔ.

	R2:	Average of non-malignant lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, a							
	Pkey	ExAcon	UnigenelD	Unigene Tille	R1	R2			
25	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	132.45	4.00			
	400666			NM_002425:Homo sapiens matrix metallopro	3.26	3.22			
	401780			NM_005557*:Homo sapiens keratin 16 (foca	26.47	10.50			
	401781			Target Exon	10.33	4.61			
	401785			NM_002275*:Homo sapiens keratin 15 (KRT1	4.13	2.70			
30	401994	٠.		Target Exon	61.84	47.00			
	402075			ENSP00000251056*:Plasma membrane calcium	1.00	1.00			
	404996			Target Exon	1.00	1.00			
	407839	AA045144	Hs.161566	ESTs	173.91	108.00			
~ -	408000	L11690	Hs.620	bullous pemphigoid antigen 1 (230/240kD)	151.17	8.00			
35	408522	A1541214	Hs.46320	Small proline-rich protein SPRK [human,	1.98	1.24			
	410561	BE540255	Hs.6994	Homo sapiens cDNA: FLJ22044 fis, clone H	10.04	1.00			
	415091	AL044872	Hs.77910	3-hydroxy-3-methylglutaryl-Coenzyme A sy	1.00	30.00			
	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t	24.30	1.00			
40	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara	53.29	51.00			
40	417034	NM_006183	Hs.80962	neurotensin	1.00	1.00			
	417366	BE185289	Hs.1076	small proline-rich protein 1B (comifin)	8.97	3.27			
	418663	AK001100	Hs.41690	desmocollin 3	112.17	19.00			
	418678	NM_001327	Hs.87225	cancer/testis antigen	1.18	1.10			
45	419121	AA374372	Hs.89626	parathyroid hormone-like hormone	1.00	1.00			
45	420783	AI659838	Hs.99923	lectin, galactoside-binding, soluble, 7	3.04	1.25			
	421773	W69233	Hs.112457	ESTs	1.12	1.14			
	421948	L42583	Hs.334309	keratin 6A	51.83	20.25			
	421978	AJ243662	Hs.110196	NICE-1 protein	1.01	0.91			
50	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	2.37	1.10			
50	422440	NM_004812	Hs.116724	aldo-kelo reductase family 1, member B10	47.53	32.00			
	423634	AW959908	Hs.1690	heparin-binding growth factor binding pr	76.02	1.00			
	423725	AJ403108	Hs.132127	hypothetical protein LOC57822	4.20	1.00			
	423738	AB002134	Hs.132195	airway trypsin-like protease	10.14	51.00			
55	424012	AW368377	Hs.137569	turnor protein 63 kDa with strong homolog	233.42	68.00			
23	424046	AF027866	Hs.138202	serine (or cysteine) proteinase inhibito	1.00	1.00			
	424098	AF077374	Hs.139322	small proline-rich protein 3	137.82	54.00			
	424834	AK001432	Hs.153408	Homo sapiens cDNA FLJ10570 fis, clone NT	56.19 . 33.45	12.00 1.00			
	425650	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	4.24	17.00			
60	427099	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophlia, mous	51.83	4.00			
UU	427335	AA448542	Hs.251677	G antigen 7B	1.00	1.00			
	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT	1.00	16.00			
	428645	AA431400	Hs.98729	ESTs, Weakly similar to 2017205A dihydro Ksp37 protein	1.00	87.00			
	428748	AW593206	Hs.98785 Hs.292911	ESTs, Highly similar to S60712 band-6-pr	2.01	1.18			
65	429259 429538	AA420450 BE182592	Hs.11261	small profine-rich protein 2A	4.43	2.90			
05	429338 429903	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	11.80	1.00			
	429903	BE062109	Hs.241551	chloride channel, calcium activated, fam	12.28	41.00			
	430480	X54232	Hs.2699	glypican 1	1.58	1.40			
	431009	BE149762	Hs.48956	gap junction protein, beta 6 (connexin 3	60.25	28.00			
70	431846	BE019924	Hs.271580	uroplakin 1B	4.49	2.51			
,,	433091	Y12642	Hs.3185	lymphocyte antigen 6 complex, locus D	1.20	1.09			
	434360	AW015415	Hs.127780	ESTs	40.98	27.00			
	434880	U02388	Hs.101	cytochrome P450, subfamily IVF, polypept	1.00	1.00			
	435505	AF200492	Hs.211238	interleukin-1 homolog 1	1.00	38.00			
75	435793	AB037734	Hs.4993	KIAA1313 protein	23.68	42.00			
, 5	436511	AA721252	Hs.291502	ESTs	16.76	14.00			
	438403	AA806607	Hs.292206	ESTs	1.00	1.00			
	439285	AL133916	113.232200	hypothetical protein FLJ20093	46.23	139.00			
	439606	W79123	Hs.58561	G protein-coupled receptor 87	33.61	1.00			
80	439670	AF088076	Hs.59507	ESTs, Weakly similar to AC004858 3 U1 sm	1.00	1.00			
	439706	AW872527	Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH	86.55	11.00			
	440325	NM_003812	Hs.7164	a disintegrin and metalloproteinase doma	62.88	147.00			
	441525	AW241867	Hs.127728	ESTs	1.53	1.42			
	443162	T49951	Hs.9029	DKFZP434G032 protein	31.11	38.00			
85	444378	R41339	Hs.12569	ESTs	1.00	1.00			
					_				

	W	O 02/086	443				PCT/US02/12476				
5	446292 447078 447342 449003 449101 450832 452240 453317 453830 454098 455601	AF081497 AW885727 A1199268 X76342 AA205847 AW970602 AI591147 NM_002277 AA534296 W27953 AI368680	Hs.279682 Hs.9914 Hs.19322 Hs.389 Hs.23016 Hs.105421 Hs.61232 Hs.41696 Hs.20953 Hs.292911 Hs.816	Rh type C glycoprotein ESTs Homo sapiens, Similar to RiKEN cDNA 2010 alcohol dehydrogenase 7 (class IV), mu o G protein-coupled receptor ESTs ESTs keratin, hair, acidic, 1 ESTs ESTs, Highly similar to S60712 band-6-pr SRY (sex determining region Y)-box 2	1.55 47.24 28.63 1.00 2.58 25.17 13.42 1.19 24.92 1.26 206.11	1.26 24.00 1.00 1.00 27.00 36.00 1.00 1.27 25.00 1.11	·				
	TABLE 12E	1									
15	Pkey: CAT number Accession:	er: Gene cluste	probeset iden er number ecession numb								
20	Pkey 439285	CAT Number 47065_1	AL13391	on 16 N79113 AF086101 N76721 AW950828 AA36401: 52 N62351 N59253 AA626243 Al341407 BE175639	3 AW955684 Al34 AA456968 Al3589	6341 A1867454 N54784 / 918 AA457077	A1655270 A1421279 AW014882				
25	TABLE 120	TABLE 12C .									
23	Pkey: Ref:	Ref: Sequence source. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The UNA									
30	Strand: Indicates DNA strand from which exons were predicted. Nt_position: Indicates nucleotide positions of predicted exons.										
35	Pkey 400666 401780 401781 401785 401994 402075 404996	Ref 8118496 7249190 7249190 7249190 4153858 8117407 6007890	Strand Plus Minus Minus Minus Minus Minus Plus Plus	Nl_position 17982-18115,20297-20456 28397-28617,28920-29045,29135-29296,29411-2 83215-83435,83531-83656,83740-83901,84237-8 165776-165996,166189-166314,166408-166569, 42904-43124,43211-43336,44607-44763,45199-4 121907-122035,122804-122921,124019-124161, 37999-38145,38652-38998,39727-39872,40557-4	4393,84955-8503 167112-167268,16 5281,46337-4673 124455-124610,12	7,86290-86814 17387-167469,168634-16 2 15672-126076	8942				
40											

TABLE 13A: Genes Distinguishing Non-Malignant Lung Disease from Lung Tumors and Normal lung

5

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Pkey

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402075

Ref

8117407

Strand

. Nt_position

Table 13A shows about 23 genes upregulated in non-malignant lung disease relative to lung tumors and normal lung. These genes were selected from about 59680 probesets on the Eos/Affyrmetrix Hu03 Genechip array.

Table 13B show the accession numbers for those Pkey's lacking UnigenelD's for table 13A. For each probeset we have listed the gene cluster number from which the oligonucleolides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 13C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 13A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

						•
15	Pkey:		et identifier number			
	ExAcon:	Exemplar Accession	n number, Genbank accession number			
	UnigenelD:	Unigene number				
	Unigene Title:	Unigene gene title				
	R1:		nors (including squamous cell carcinomas, adenocar	cinomas, small cell c	carcinomas, granulomatous and	carcinoid turnors) divided by the
20		average of normal	lung samples			
	R2:	Average of non-ma	lignant lung disease samples (including bronchitis, e	nphysema, fibrosis, a	atelectasis, asthma) divided by t	he average of normal lung samples
	Pkey ExAc		Unigene Title	R1	R2	
25	408562 Al436		Homo sapiens mRNA for KIAA1568 protein,	1.00	230.00	
25	409031 AA37		ESTs	1.00	128.00	
	412372 R659		hypothetical protein FLJ22029	1.00	173.00	
	415910 U203		chemokine (C-X3-C) receptor 1	1.00	145.00	
	417511 AL04		chordin-like	1.00	179.00	
20	418819 AA22		ESTs	1.00	140.00	
30	422060 R208		ESTs, Moderately similar to ALU5_HUMAN A	1.00	156.00	
	424585 AA46		ESTs	1.00	167.00	
	426753 T898	32 Hs.170278	ESTs	1.00	141.00	
	429496 AA45		ESTs	1.00	138.00	
25	430719 AA48		ESTs	1.00	133.00	
35	431089 BE04		ESTs, Weakly similar to unknown protein	23.32	941.00 157.00	
	431385 BE17		membrane-spanning 4-domains, subfamily A	1.00	157.00	
		07351 Hs.268107	multimerin	1.00 1.00	218.00	
	436532 AA72		gb:nv54h12.r1 NCI_CGAP_Ew1 Homo sapiens	1.00	147.00	
40	437960 Al669		ESTs	1.00	141.00	
40	438202 AW16		ESTs	1.00	167.00	
	441499 AW29		ESTs glutamate receptor, ionotropic, AMPA 1	1.00	151.00	
	444513 AL126			1.00	141.00	•
	448253 H258 453636 R678		ESTs ESTs	1.00	116.00	
45	458332 Al000		ESTs	.1.00	192.00	
73	459587 AA03		gb:zk15e04.s1 Soares_pregnant_uterus_NbH	1.00	154.00	٠.
	409001 AAUS	1930	gu.zk13804.51 30ales_pregnant_uterus_Ruri	1.00	104.00	
	TABLE 13B				·	
	TAULE 100					
50	Pkey: U	nique Eos probeset is	tentifier number			
-		ene cluster number				
		enbank accession nu	mbers		•	
	,		,			
	Pkey C	AT Number Accession	on			
55	431089 32	27825_1 BE04139	95 AA491826 AA621946 AA715980 AA666102		•	
	436532 42	21802_1 AA7215	22 AW975443 T93070			
		_				
			, .	•		
~	TABLE 13C	•				
60		. •		•		
	Pkey: U	nique number corres	ponding to an Eos probeset			H
	Ref: Se	equence source. The	7 digit numbers in this column are Genbank Identific	er (GI) numbers. "Du	unnam i. et al." refers to the publ	cauon entitieo "The DNA
	ŞE	equence of human ch	romosome 22." Dunham I. et al., Nature (1999) 402	489-495.		
65			from which exons were predicted.	•		
65	Nt_position: In	dicates nucleotide po	sitions of predicted exons.			

121907-122035,122804-122921,124019-124161,124455-124610,125672-126076

Table 14A shows the subcellular localization and preferred utility for the genes appearing in Tables 9A and 10A. mAb symbolizes monoclonal antibody, diag symbolizes diagnostic, s.m. symbolizes small molecule, and CTL symbolizes cytotoxic lymphocytic ligand. These genes were selected from 59680 probesets on the Eos/Afflymetrix Hu03 5 Genechip array.

Table 14B show the accession numbers for those Pkey's tacking UnigenelD's for table 14A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 14C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 14A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

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Pkey: ExAcon:

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number

Example Accession number Unigene Title: Unigene Title: Unigene gene title
Pref. Utility: Preferred Utility
Pred. Loc: Predicted subcellular localization

	rieu.Luc	. Freuitieu a	Subcellulai ibtali	280011		
	Pkey	ExAcon	UnigenelD	Unigene Title	Pref Utility	Pred. Loc
	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	mAb & diag & s.m.	extracellular
25	400303	AA242758	Hs.79136	LIV-1 protein, estrogen regulated	mAb	plasma membrane
	402075	701242100	110.10100	ENSP00000251056*:Plasma membrane caldium		secreted
	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon	diag	secreted
	408243	Y00787	Hs.624	interleukin 8	diag	secreted
	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,	mAb & s.m.	plasma membrane
30	408908	BE296227	Hs.250822	serine/threonine kinase 15	s.m.	cytoplasm
20	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119	CTL & diag	secreted
	409103	AF251237	Hs.112208	XAGE-1 protein	CTL	nuclear
	409420	Z15008	Hs.54451	taminin, gamma 2 (nicein (100kD), kalini	diag	secreted
	409632	W74001	Hs.55279	serine (or cysteine) proteinase inhibito	diag	secreted
35	409757	NM_001898	Hs.123114	cystatin SN	diag	extracellular
55	409893	AW247090	Hs.57101	minichromosome maintenance deficient (S.	CTL	nuclear
	409956	AW103364	Hs.727	Inhibin, beta A (activin A, activin AB a	diag	extracellular
	410001	AB041036	Hs.57771	kalikrein 11	dian	extracellular
	410407	X66839	Hs.63287	carbonic anhydrase IX	mAb & s.m.	plasma membrane
40	410418	D31382	Hs.63325	transmembrane protease, serine 4	mAb & diag & s.m.	plasma membrane
-70	412140	AA219691	Hs.73625	RAB6 Interacting, kinesin-like (rabkines	s.m.	picerile manierano
	412719	AW016610	Hs.816	ESTs	s.m.	nuclear
	414774	X02419	Hs.77274	plasminogen activator, urokinase	diag	extracellular
	414883	AA926960	16.77274	CDC28 protein kinase 1	s.m.	310 00011010
45	415138	C18356	Hs.295944	tissue factor pathway inhibitor 2	CTL & diag	extracellular
15	415669	NM_005025	Hs.78589	serine (or cystelne) proteinase inhibito	mAb & diag & s.m.	secreted
	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t	mAb & s.m.	plasma membrane
	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara	diag	extracellular
	417034	NM_006183	Hs.80962	neurotensin	diag	extracellular
50	417079	U65590	Hs.81134	interleukin 1 receptor antagonist	diag	extracellular
50	417308	H60720	Hs.81892	KIAA0101 gene product	s.m.	mitochondrial
	417389	BE260964	Hs.82045	midkine (neurite growth-promoting factor	mAb & dlag	secreted
	417433	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein	mAb	plasma membrane
	417933	X02308	Hs.82962	thymidylate synthetase	s.m.	endoplasmic reticulum
55	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me	s.m.	cytoplasm
	418506	AA084248	Hs.85339	G protein-coupled receptor 39	mAb & s.m.	plasma membrane
	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)	CTL	cytoplasmic
	419121	AA374372	Hs.89626	parathyroid hormone-like hormone	diag	secreted
	419171	NM_002846	Hs.89655	protein tyrosine phosphatase, receptor t	mAb & s.m.	plasma membrane
60	419183	U60669	Hs.89663	cytochrome P450, subfamily XXIV (vitamin	CTL & s.m.	mitochendrial
	419216	AU076718	Hs.164021	small inducible cytokine subfamily B (Cy	diag	secreted
	419235	AW470411	Hs.288433	neurotrimin	mAb & diag	plasma membrane
	419452	U33635	Hs.90572	PTK7 protein tyrosine kinase 7	mAb & s.m.	plasma membrane
	419556	U29615	Hs.91093	chitinase 1 (chitotriosidase)	mAb & diag	extracellular*
65	420610	Al683183	Hs.99348	distal-less homeo box 5	CTL	nuclear
•	421110	AJ250717	Hs.1355	cathepsin E	sm & diag	extracellular
	421379	Y15221	Hs.103932	small inducible cytokine subfamily B (Cy	diag	secreted
	421474	U76362	Hs.104637	solute carrier family 1 (glutamate trans	mAb & s.m.	plasma membrane
	421552	AF026692	Hs.105700	secreted frizzled-related protein 4	diag	secreted
70	421753	BE314828	Hs.107911	ATP-binding cassette, sub-family B (MDR/	mAb & s.m.	plasma membrane
	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	mAb & s.m.	plasma membrane
	422109	S73265	Hs.1473	gastrin-releasing peptide	diag	secreted
	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	diag	secreted
	422282	AF019225	Hs.114309	apolipoprotein L	diag	secreted
<i>75</i>	422283	AW411307	Hs.114311	CDC45 (cell division cycle 45, S.cerevis	s.m.	nuclear
	422424	Al186431	Hs.296638	prostate differentiation factor	diag	extracellular
•	422765	AW409701	Hs.1578	baculoviral IAP repeat-containing 5 (sur	s.m.	cyloplasm
	422809	AK001379	Hs.121028	hypothetical protein FLJ 10549	s.m.	nuclear
00	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	diag	extracellular
80	422956	BE545072	Hs.122579	ECT2 protein (Epithelial cell transformi	CTL & s.m.	
	423634	AW959908	Hs.1690	heparin-binding growth factor binding pr	diag	
	423573	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage	mAb & diag & s.m.	secreted
	423961	D13666	Hs.136348	periostin (OSF-2os)	mAb & diag	extracellular
0.5	424046	AF027866	Hs.138202	serine (or cystelne) proteinase inhibito	dlag	secreted
85	424381	AA285249	Hs.146329	protein kinase Chk2	s.m.	nuclear

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	w	O 02/086	443			
	424502	AF242388	Hs.149585	lengsin	s.m.	cytoplasmic
	424503	NM_002205	Hs.149609	integrin, alpha 5 (fibronectin receptor,	mAb & s.m.	plasma membrane
	424687	J05070	Hs.151738	matrix metalloproteinase 9 (gelatinase B	diag	extracellular
5	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin	mAb & diag & s.m.	secreted cytoplasmic
,	425322 425650	U63630 NM_001944	Hs.155637 Hs.1925	protein kinase, DNA-activated, catalytic desmoglein 3 (pemphigus vulgaris antigen	s.m. mAb	plasma membrane
	425734	AF056209	Hs.159396	peptidylglycine alpha-amidating monooxyg	s.m.	F 1
	425776	U25128	Hs.159499	parathyroid hormone receptor 2	mAb & diag	plasma membrane
10	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily member	mAb & s.m.	plasma membrane
10	426215 426427	AW963419 M86699	Hs.155223 Hs.169840	stanniocaldin 2 TTK protein kinase	mAb & diag CTL & s.m.	secreted nuclear
	426514	BE616633	Hs.170195	bone morphogenetic protein 7 (osteogenic	mAb & diag	secreted
	427335	AA448542	Hs.251677	G antigen 7B	CTL	cytoplasmic
1.5	427747	AW411425	Hs.180655	serine/threonine kinase 12	s.m.	cytoplasmic
15	428242	H55709	Hs.2250	leukemia inhibitory factor (cholinergic	diag	extracellular
	428330 428450	L22524 NM_014791	Hs.2256 Hs.184339	matrix metalioproteinase 7 (matrilysin, KIAA0175 gene product	mAb & diag & s.m. s.m.	nuclear
	428479	Y00272	Hs.334562	cell division cycle 2, G1 to S and G2 to	s.m.	uncjest
	428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino	mAb & s.m.	plasma membrane
20	428664	AK001666	Hs.189095	similar to SALL1 (sal (Drosophila)-like	CTL & s.m.	nuclear
	428698	AA852773	Hs.334838	KIAA1866 protein	mAb	extracellular
	428748 428758	AW593206 AA433988	Hs.98785 Hs.98502	Ksp37 protein CA125 antigen; mucin 16	diag diag	mitochodria*
	428969	AF120274	Hs.194689	artemin	diag	extracellular
25	429211	AF052693	Hs.198249	gap junction protein, beta 5 (connexin 3	mAb & s.m.	plasma membrane
	429263	AA019004	Hs.198396	ATP-binding cassette, sub-family A (ABC1	mAb & s.m.	plasma membrane
	429547	AW009166	Hs.99376	ESTS	diag	secreted secreted
	429610 429903	AB024937 AL134197	Hs.211092 Hs.93597	LUNX protein; PLUNC (palate lung and nas cyclin-dependent kinase 5, regulatory su	mAb & diag s.m.	Secreted
30	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam	mAb & s.m.	plasma membrane
	431462	AW583672	Hs.256311	granin-like neuroendocrine peptide precu	diag	extracellular
	431515	NM_012152	Hs.258583	endothelial differentiation, lysophospha	mAb & s.m.	plasma membrane
	431846	BE019924	Hs.271580	uroplakin 1B	mAb & diag mAb & diag	plasma membrane plasma membrane
35	431958 432201	X63629 AI538613	Hs.2877 Hs.298241	cadherin 3, type 1, P-cadherin (placenta Transmembrane protease, serine 3	mAb & diag & s.m.	plasma membrane
55	433001	AF217513	Hs.279905	clone HQ0310 PRO0310p1	s.m.	nuclear
	435505	AF200492	Hs.211238	interleukin-1 homolog 1	diag	secreted
	436481	AA379597	Hs.5199	HSPC150 protein similar to ubiquitin-con	s.m.	adonlarm
40	437016 437044	AU076916 AL035864	Hs.5398 Hs.69517	guanine monphosphate synthetase differentially expressed in Fanconi's an	s.m. CTL	cytoplasm ER
70	437789	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti	CTL	nuclear
	437852	BE001836	Hs.256897	ESTs, Weakly similar to dJ365O12.1 [H.sa	mAb & s.m.	plasma membrane
	439223	AW238299	Hs.250618	UL16 binding protein 2	mAb	plasma membrane
45	439477 439606	W69813	Hs.58042 Hs.58561	ESTs, Moderately similar to GFR3_HUMAN G	mAb & s.m. mAb & s.m.	plasma membrane
7.7	439738	W79123 BE246502	Hs.9598	G protein-coupled receptor 87 sema domain, immunoglobulin domain (Ig),	mAb & s.m.	plasma membrane
	440006	AK000517	Hs.6844	NALP2 protein; PYRIN-Containing APAF1-II	s.m.	nuclear
	441362	BE614410	Hs.23044	RAD51 (S. cerevisiae) homolog (E coli Re	s.m.	
50	442117	AW664964	Hs.128899	ESTs; hypothetical protein for IMAGE:447	mAb & s.m.	plasma membrane extracellular
30	443247 443426	BE614387 AF098158	Hs.333893 Hs.9329	c-Myc target JPO1 chromosome 20 open reading frame 1	CTL CTL	extracential
	443859	NM_013409	Hs.9914	follistatin	diag	extracellular
	444006	BE395085	Hs.10086	type I transmembrane protein Fn14	mAb	plasma membrane
55	444371	BE540274	Hs.239	forkhead box M1	s.m.	nuclear
33	444381 444781	BE387335 NM_014400	Hs.283713 Hs.11950	ESTs, Weakly similar to S64054 hypotheti GPI-anchored metastasis-associated prote	diag mAb & diag	secreted plasma membrane
	445537	AJ245671	Hs.12844	EGF-like-domain, multiple 6	mAb & diag	secreted
	446619	AU076643	Hs.313	secreted phosphoprotein 1 (osteopontin,	diag	secreted
60	446921	AB012113	Hs.16530	small inducible cytokine subfamily A (Cy	diag	extracellular
60	447033	AI357412	Hs.157601	ESTS	CTL & diag	secreted
	447342 448243	AI199268 AW369771	Hs.19322 Hs.52620	Homo saplens, Similar to RIKEN cDNA 2010 integrin, beta 8	CTL mAb & s.m	plasma membrane
	448844	Al581519	Hs.177164	ESTs -	mAb & s.m.	processor mornor and
	449048	Z45051	Hs.22920	similar to S68401 (cattle) glucose induc	πAb	plasma membrana
65	449722	BE280074	Hs.23960	cyclin B1	s.m.	cytoplasm
	450001	NM_001044	Hs.406	solute carrier family 6 (neurotransmitte	mAb & s.m.	plasma membrane plasma membrane
	450375 450701	AA009647 H39960	Hs.288467	a disintegrin and metalloproteinase doma hypothetical protein XP_098151 (leucine-	mAb & diag & s.m. mAb & diag	plasma membrane
	450983	AA305384	Hs.25740	ERO1 (S. cerevisiae)-like	diag	secreted
70	451668	Z43948	Hs.326444	cartilage acidic protein 1	mAb & diag	plasma membrane
	452281	T93500	Hs.28792	Homo sapiens cDNA FLJ11041 fis, clone PL	diag	autona illutar
	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro Ig superfamily receptor LNIR	diag mAb	extracellular plasma membrane
	452747 452838	BE153855 U65011	Hs.61460 Hs.30743	preferentially expressed antigen in mela	CTL	nuclear
75	453968	AA847843	Hs.62711	High mobility group (nonhistone chromoso	CTL & s.m.	nuclear
	457489	A1693B15	Hs.127179	cryptic gene	diag	secreted
	TABLE 1	4B				

TABLE 14B

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Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers Pkey CAT Number Accession

	wo	02/08644	3	•	PCT/US02/12476
5	414883	15024_1	AA08243 AA29275 AA87203 R75953 A AW61300 N95210 A	0 AA926959 W76521 W24270 W21526 AA037172 BE267636 H83186 AA469909 N86396 AA001346 6 H72525 H77575 N49786 W80565 H78746 BE569085 W04339 R98127 T55938 BE279271 AW960 3 AA177048 NM_001826 X54941 BE314366 AA908783 AI719075 BE270172 BE269819 AA889955 9 W72395 T99630 Al422691 H98460 N31428 BE255916 H03265 AI857576 AA776920 AA910644 A \W662396 AA662522 AI865147 AI423153 AW262230 AA584410 AA583187 AW024595 AW069734 \U20122 AA527373 AW972459 AI831360 AA621337 AA10926 AA772418 AA59628 AI033892 W95096 A \U2014459432 AI041437 AA932124 AA627684 AA935829 AI004827 AI423513 AI094597 H42079 R54703 \U2015 W44561 AI991988 AI537692 AI090262 AA740817 AI312104 AI911822 AA416871 AI185409 AA12	1304 T29812 AA476873 BE297387 A1204630 W25243 A1935150 A459522 AA293140 AW514667 A1828996 AA282997 AA876046 A1034317 AA398727 A1085031 I A1630359 AA617681 AA978045
10			Al139549 Al494230 Al494211 AA95434	AA633648 AI339996 AI336880 AA399239 AI078708 AI085351 AI362835 AI346618 AI146955 AI986 AI278887 AA962596 AI492600 W80435 AA001979 R97424 AI129015 N24127 AA157451 AA23554 AW059501 AW886710 R92790 N59755 AI361128 AW589407 H47725 H97534 H48076 H484501 H77576 R96823 AI457100 N92845 N49682 H42038 BE220698 BE220715 H99552 AA701624 N74 E281919 AA769633 AA480310 AA507454 AA910586 AI203723 AW104725 W25611 W25071 T889	9380 Al348243 N92892 AA765850 19 AA459292 AA037114 AA129785 9631 AW300758 H03431 R76789 1173 R54704 H79520 H72923
15	450375	B3327_1	W95095 F AA009647	REZ01919 AA703533 AA460310 AA5011952 H82956 N83673 AA283672 97470 AA702275 T77551 AA911952 H82956 N83673 AA283672 7 AA131254 AA374293 AW954405 H04410 AW606284 AA161166 BE157467 BE157601 H04384 W 3 H03231 H59605 H01642 AA852876 AA113758 AA626915 AA746952 A1161014 AA099554 R6906	46291 AW663674 H04021 H01532
20	TABLE 14C				
20	Pkey: Ref:	Sequence sour	ce. The 7 di	ing to an Eos probeset git numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publ Isome 22." Dunham I. et al., Nature (1999) 402:489-495.	ication entitled "The DNA
25	Strand: Nt_position:	Indicates DNA	strand from v	which exons were predicted. ns of predicted exons.	
	Pkey	Ref S	Strand	Nt_position	
20	402075	8117407	Plus	121907-122035,122804-122921,124019-124161,124455-124610,125672-126076	

TABLE 15A: Information for all sequences in Table 16

Table 15A shows the Seq ID No, Pkey, ExAcon, UnigeneID, and Unigene Title for all of the sequences in Table 16.

- Table 15B show the accession numbers for those Pkey's lacking UnigenelD's for table 15A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence stmilarly using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the 5 "Accession" column.
- Table 15C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 15A. For each predicted exon, we have listed the genomic 10 sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Seq ID No: Sequence ID number Pkey: Unique Eos probeset 15

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number ExAcon:

UnigenelD: Unigene number Unigene Title: Unigene gene title

	Ottidette tinerounderte A	CITO UUB			
20	Seq ID No:	Pkey	ExAcon	UnigenelD	Unigene Title
	Seq ID No: 1 & 2	410407	X66839	Hs.63287	carbonic anhydrase IX
	Seq ID No: 3 & 4	412719	AW016610	Hs.816	ESTs
	Seq ID No: 5 & 6	417034	NM_006183	Hs.80962	neurotensin
25	Seq ID No: 7 & 8	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam
	Seq ID No: 9 & 10	407788	BE514982	Hs.38991	S100 calcium-binding protein A2
	Seq ID No: 11 & 12	407788	BE514982	Hs.38991	S100 calcium-binding protein A2
	Seq ID No: 13 & 14	407788	BE514982	Hs.38991	S100 calcium-binding protein A2
30	Seq ID No: 15 & 16	407788	BE514982	Hs.38991	S100 calcium-binding protein A2 hypothetical protein FLJ20093
30	Seq ID No: 17 & 18 Seq ID No: 19 & 20	439285 413753	AL133916 U17760	Hs.75517	Iaminin, beta 3 (nicein (125kD), kalinin
	Seq ID No: 21 & 22	120486	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog
	Seq ID No: 23 & 24		NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen
	Seq ID No: 25 & 26	412140	- AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines
35	Seq ID No: 27 & 28	423673	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage
	Seq ID No: 29 & 30	452838	U65011	Hs.30743	preferentially expressed antigen in mela
	Seq ID No: 31 & 32	418663	AK001100	Hs.41690	desmocollin 3
	Seq ID No: 33 & 34	418663	AK001100	Hs.41690	desmocollin 3
40	Seq ID No: 35 & 36	409632	W74001	Hs.55279	serine (or cysteine) proteinase inhibito LUNX protein; PLUNC (palate lung and nas
40	Seq ID No: 37 & 38	429610 406690	AB024937 M29540	Hs.211092 Hs.220529	carcinoembryonic antigen-related cell ad
	Seq ID No: 39 & 40 Seq ID No: 41 & 42	431846	BE019924	Hs.271580	uroplakin 1B
	Seq ID No: 43 & 44	418830	BE513731	Hs.88959	hypothetical protein MGC4816
	Seg ID No: 45 & 46	424098	AF077374	Hs.139322	smail proline-rich protein 3
45	Seq ID No: 47 & 48	443648	A1085377	Hs.143610	ESTs
	Seq ID No: 49	311034	BE567130	Hs.311389	ESTs, Highly similar to NKGD_HUMAN NKG2-
	Seq ID No: 50 & 51	408522	A1541214	Hs.46320	Small proline-rich protein SPRK [human,
	Seq ID No: 52 & 53	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL
50	Seq ID No: 54 & 55	435505	AF200492	Hs.211238 Hs.1076	interleukin-1 homolog 1 small proline-rich protein 1B (comifin)
30	Seq ID No: 56 & 57	417366 431958	BE185289 X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta
•	Seq ID No: 58 & 59 Seq ID No: 60 & 61	441020	W79283	Hs.35962	ESTs
	Seq ID No: 62 & 63	423217	NM 000094	Hs.1640	collagen, type VII, alpha 1 (epidermolys
	Seq ID No: 64 & 65	429538	BE182592	Hs.11261	small proline-rich protein 2A
55	Seq ID No: 66 & 67	448733	NM_005629	Hs.187958	solute carrier family 6 (neurotransmitte
	Seq ID No: 68 & 69	444371	BE540274	Hs.239	forkhead box M1
	Seq ID No: 70 & 71	444371	BE540274	Hs.239	forkhead box M1
	Seq ID No: 72 & 73	444371	BE540274	Hs:239 Hs.112408	forkhead box M1 S100 calcium-binding protein A7 (psorias
60	Seq ID No: 74 & 75 Seq ID No: 76 & 77	422168 422168	AA586894 AA586894	Hs.112408	\$100 calcium-binding protein A7 (psorias
00	Seq ID No: 78 & 79	429259	AA420450	Hs.292911	Plakophilin
	Seq ID No: 80 & 81	426440	BE382756	Hs.169902	solute carrier family 2 (facilitated glu
•	Seq ID No: 82 & 83	437044	AL035864	Hs.69517	differentially expressed in Fanconi's an
	Seq ID No: 84 & 85	423662	AK001035	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro
65	Seq ID No: 86 & 87	428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino
	Seq ID No: 88 & 89	429211	AF052693	Hs.198249	gap junction protein, beta 5 (connexin 3
	Seq ID No: 90 & 91	417389	BE260964 AW959908	Hs.82045 Hs.1690	midkine (neurite growth-promoting factor heparin-binding growth factor binding pr
	Seq ID No: 92 & 93 Seq ID No: 94 & 95	423634 417515	L24203	Hs.82237	ataxia-telangiectasia group D-associated
70	Seq ID No: 96 & 97	441362	BE614410	Hs.23044	RAD51 (S. cerevisiae) homolog (E coli Re
, •	Seq ID No: 98 & 99	425322	U63630	Hs.155637	protein kinase, DNA-activated, catalytic
	Seq ID No: 100 & 101	449003	X76342	Hs.389	alcohol dehydrogenase 7 (class IV), mu o
	Seq ID No: 102 & 103	431009	BE149762	Hs.48956	gap junction protein, beta 6 (connexin 3
75	Seq ID No: 104 & 105	409103	AF251237	Hs.112208	XAGE-1 protein
75	Seq ID No: 106 & 107	417542	J04129	Hs.82269	progestagen-associated endometrial prote stratifin
	Seq ID No: 108 & 109	428471	X57348 U37519	Hs.184510 Hs.87539	aldehyde dehydrogenase 3 family, member
	Seq ID No: 110 & 111 Seq ID No: 112 & 113	418004 414761	AU077228	Hs.77256	enhancer of zeste (Drosophila) homolog 2
	Seq ID No: 114 & 115	418203	X54942	Hs.83758	CDC28 protein kinase 2
80	Seq ID No: 116	447343	AA256841	Hs.236894	ESTs, Highly similar to S02392 alpha-2-m
	Seq ID No: 117 & 118	437016	AU076916	Hs.5398	guanine monphosphate synthetase
	Seq ID No: 119 & 120	449230	BE613348	Hs.211579	melanoma cell adhesion molecule
	Seq ID No: 121 & 122	446989	AK001898	Hs.16740	hypothetical protein FLJ11036
95	Seq ID No: 123 & 124	457819	AA057484	Hs.35406	ESTs, Highly similar to unnamed protein
85	Seq ID No: 125 & 126	424687	J05070	Hs.151738	matrix metalioproteinase 9 (gelatinase B

	WO 02/086	443			
	Seq ID No: 127 & 128	414430	Al346201	Hs.76118	ubiquilin carboxyl-terminal esterase L1
	Seq ID No: 129 & 130	418462	BE001596	Hs.85266	integrin, beta 4
	Seq ID No: 131 & 132	100668	L05424	Hs.169610	CD44 antigen (homing function and Indian
~	Seq ID No: 133 & 134	458933	AI638429	Hs.24763	RAN binding protein 1
5	Seq ID No: 135 & 136	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
	Seq ID No: 137 & 138	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
	Seq ID No: 139 & 140	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
	Seq ID No: 141 & 142	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
10	Seq ID No: 143 & 144	446269	AW263155	Hs.14559	hypothetical protein FLJ10540
10	Seq ID No: 145 & 146	422765	AW409701	Hs.1578	bacutoviral IAP repeat-containing 5 (sur
	Seq ID No: 147 & 148	436481	AA379597	Hs.5199	HSPC150 protein similar to ubiquitin-con
	Seq ID No: 149 & 150	440325	NM_003812	Hs.7164	a disintegrin and metalloproteinase doma
	Seq ID No: 151 & 152	439606	W79123	Hs.58561	G protein-coupled receptor 87
1.5	Seq ID No: 153 & 154	453884	AA355925	Hs.36232	KIAA0186 gene product
15	Seq ID No: 155 & 156	453884	AA355925	Hs.36232	KIAA0186 gene product
	Seq ID No: 157 & 158	453884	AA355925	Hs.36232	KIAA0186 gene product
	Seq ID No: 159 & 160	453884	AA355925	Hs.36232	KIAA0186 gene product
	Seq ID No: 161 & 162	404877			NM_005365:Homo sapiens melanoma antigen,
20	Seq ID No: 163 & 164	413129	AF292100	Hs.104613	RP42 homolog
20	Seq ID No: 165 & 166	413281	AA851271	Hs.222024	transcription factor BMAL2
	Seq ID No: 167 & 168	444781	NM_014400	Hs.11950	GPI-anchored metastasis-associated prote
	Seq ID No: 169 & 170	416819	U77735	Hs.80205	pim-2 oncogene
	Seq ID No: 171 & 172	451320	AW118072		diacylglycerol kinase, zeta (104kD)
0.5	Seq ID No: 173 & 174	418543	NM_005329	Hs.85962	hyaluronan synthase 3
25	Seq ID No: 175 & 176	454034	NM_000691	Hs.575	aldehyde dehydrogen ase 3 family, member
	Seq ID No: 177 & 178	425397	J04088	Hs.156346	topoisomerase (DNA) II alpha (170kD)
	Seq ID No: 179 & 180	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 181 & 182	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 183 & 184	415817	U88967	Hs.78867	protein tyrosine phosphalase, receptor-t
30	Seq ID No: 185 & 186	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 187 & 188	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
٩	Seq ID No: 189 & 190	419121	AA374372	Hs.89626	parathyrold hormone-like hormone
	Seq ID No: 191 & 192	448993	AI471630	Hs.8127	KIAA0144 gene product
~ ~	Seq ID No: 193 & 194	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR
35	Seq ID No: 195 & 196	430393	BE185030	Hs.241305	estrogen-responsive B box protein
	Seq ID No: 197 & 198	425057	AA826434	Hs.1619	achaete-scute complex (Drosophila) homol
	Seq ID No: 199 & 200	420462	AF050147	Hs.97932	chondromodulin I precursor
	Seq ID No: 201 & 202	102963	X02404	Hs.274534	calcitonin-related polypeptide, beta
40	Seq ID No: 203 & 204	100576	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid
40	Seq ID No: 205 & 206	101175	U82671	Hs.36980	melanoma antigen, family A, 2
	Seq ID No: 207 & 208	429038	AL023513	Hs.194766	seizure related gene 6 (mouse)-like
	Seq ID No: 209 & 210	418678	NM_001327	Hs.167379	cancer/testls antigen (NY-ESO-1)
	Seq ID No: 211 & 212	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
45	Seq ID No: 213 & 214	131927	AJ003112	Hs.34780	doublecortex; lissencephaly, X-linked (d
45	Seq ID No: 215 & 216	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT
	Seq ID No: 217 & 218	427335	AA448542	Hs.251677	G antigen 7B
		409420	Z15008	Hs.54451	laminin, gamma 2 (nicein (100kD), kalini
	Seq ID No: 219 & 220		** *****		
	Seq ID No: 221 & 222	114346	AL137256	Hs.130489	ATPase, aminophospholipid transporter-li
50	Seq ID No: 221 & 222 Seq ID No: 223 & 224	114346 438956	AL137256 W00847	Hs.135056	Human DNA sequence from clone RP5-850E9
50	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226	114346 438956 404440	W00847	Hs.135056	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen,
50	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228	114346 438956 404440 415669	W00847 NM_005025	Hs.135056 Hs.78589	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito
50	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 229 & 230	114346 438956 404440 415669 103312	W00847 NM_005025 Y12642	Hs.135056 Hs.78589 Hs.3185	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal
50	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 229 & 230 Seq ID No: 231 & 232	114346 438956 404440 415669 103312 320843	W00847 NM_005025 Y12642 BE069288	Hs.135056 Hs.78589 Hs.3185 Hs.34744	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr
	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 229 & 230 Seq ID No: 231 & 232 Seq ID No: 233	114346 438956 404440 415669 103312 320843 429065	W00847 NM_005025 Y12642 BE069288 Al753247	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643	Human DNA sequence from clone RP5-850E9 NM_021048:Horno sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT
50 55	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 229 & 230 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 234 & 235	114346 438956 404440 415669 103312 320843 429065 446102	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito hysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs
	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 230 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 236 & 237	114346 438956 404440 415669 103312 320843 429065 446102 330495	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleotide binding protein (G pr
	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 230 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 234 & 235 Seq ID No: 236 & 237 Seq ID No: 238	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573	W00847 NM_005025 Y12642 BE069288 A1753247 AW168067 U47924 A1733859	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleotide binding protein (G pr ESTs
	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 229 & 230 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 234 & 235 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 239 & 240	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573 428479	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to
55	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 239 & 240 Seq ID No: 241 & 242	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573 428479 428479	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2, G1 to S and G2 to
	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573 428479 428479 332180	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito hysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2, G1 to S and G2 to claudin 1
55	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 230 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 243 & 244	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573 428479 428479 332180 437915	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.3327 Hs.202312	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca
55	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 233 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 239 & 240 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 246 & 247	114346 438956 404440 415669 103312 320843 429065 446102 330495 413573 428479 428479 332180 437915 441553	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.7327 Hs.202312 Hs.121296	Human DNA sequence from clone RP5-850E9 NM_0210481Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs
55	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 223 & 228 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 233 Seq ID No: 233 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 248 & 240 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 & 247 Seq ID No: 248 & 247 Seq ID No: 248 & 247 Seq ID No: 248 & 249	114346 438956 40440 415669 103312 320843 429065 446102 330495 413573 428479 32180 437915 441553 331692	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.202312 Hs.152213	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site famil
55 60	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 246 & 247 Seq ID No: 248 & 249 Seq ID No: 248 & 249 Seq ID No: 248 & 249 Seq ID No: 250 & 251	114346 438956 40440 415669 103312 320843 429065 413573 428479 332180 437915 441553 331692 429413	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877	Human DNA sequence from clone RP5-850E9 NM_021048:Horno sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guantine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2, G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein
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55 60	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 246 & 247 Seq ID No: 246 & 247 Seq ID No: 246 & 247 Seq ID No: 250 & 251 Seq ID No: 252 & 253 Seq ID No: 254 & 255	114346 438956 40440 415669 103312 320843 429665 446102 330495 413573 428479 428479 427915 441553 331692 42943 42943 42943 42943 42943 4494	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.108923	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleotide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family
55 60	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 223 & 228 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 238 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 246 & 247 Seq ID No: 250 & 251 Seq ID No: 252 & 253 Seq ID No: 254 & 255 Seq ID No: 254 & 255 Seq ID No: 254 & 255 Seq ID No: 256 & 257	114346 438956 40440 415669 103312 320843 429065 446102 330495 413573 428479 428479 332180 437915 441553 331692 429413 4229413 4229413 448357 446292	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682	Human DNA sequence from clone RP5-850E9 NM_0210481+orno sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guantine nucleotide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein
55 60	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 256 & 257 Seq ID No: 256 & 255 Seq ID No: 256 & 255 Seq ID No: 256 & 257 Seq ID No: 256 & 255	114346 438956 40440 415669 103312 320843 429055 413573 428479 332180 437915 441553 331692 429413 422283 44832 448292 416209	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.7327 Hs.202312 Hs.202312 Hs.202312 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682 Hs.79078	Human DNA sequence from clone RP5-850E9 NM_021048:Horno sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (mitotic arrest deficient, yeast, h
556065	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 250 & 251 Seq ID No: 250 & 251 Seq ID No: 256 & 253 Seq ID No: 256 & 255 Seq ID No: 258 & 259	114346 438956 40446 415669 103312 320843 42905 4330495 413573 428479 332180 437915 441553 331692 429413 422283 448357 446209 453922	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF053306	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682 Hs.79078 Hs.36708	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to cell division cycle 2, G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (mitotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1
55 60	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 & 240 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 243 & 244 Seq ID No: 243 & 245 Seq ID No: 245 & 255 Seq ID No: 256 & 257 Seq ID No: 256 & 257 Seq ID No: 258 & 259	114346 438956 40440 415669 103312 320843 429065 446102 330495 413573 428479 428479 428479 42153 33169 429243 448357 446292 416292 416292 424046	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF083306 AF027866	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682 Hs.79078 Hs.36708 Hs.138202	Human DNA sequence from clone RP5-850E9 NM_0210481-Horno sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleotide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (mitotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1 serine (or cysteine) proteinase inhibito
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5560657075	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 250 & 251 Seq ID No: 252 & 253 Seq ID No: 252 & 253 Seq ID No: 258 & 259 Seq ID No: 258 & 259 Seq ID No: 266 & 267 Seq ID No: 268 & 269 Seq ID No: 270 & 271 Seq ID No: 274 & 275 Seq ID No: 274 & 275 Seq ID No: 276 & 277 Seq ID No: 276 & 277 Seq ID No: 278 & 279 Seq ID No: 278 & 279 Seq ID No: 282 Seq ID No: 288	114346 438956 40446 415669 103312 320843 42905 413573 428479 332180 437915 44153 331692 429413 422283 44837 446292 446292 446292 453922 424046 439223 409757 411089 428969	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF053306 AF027866 AW238299 AI553633 NM_001898 AA456454 AA721252 AF120274	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682 Hs.36708 Hs.138202 Hs.250618 Hs.326447 Hs.123114 Hs.214291 Hs.291502 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689	Human DNA sequence from clone RP5-850E9 NM_021048:Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (mitotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1 serine (or cysteine) proteinase inhibito UL16 binding protein 2 ESTs cystatin SN cell division cycle 2-like 1 (PITSLRE pr ESTs artemin artemin artemin gbye53h05.s1 Soares fetal liver spleen hypothetical protein AF301222
5560657075	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 250 & 251 Seq ID No: 252 & 253 Seq ID No: 254 & 255 Seq ID No: 256 & 257 Seq ID No: 266 & 267 Seq ID No: 268 & 269 Seq ID No: 270 & 271 Seq ID No: 276 & 277 Seq ID No: 278 & 279 Seq ID No: 278 & 279 Seq ID No: 282 & 281 Seq ID No: 283 & 284	114346 438956 40446 415669 103312 320843 42905 413573 428479 332180 437915 441553 331692 429413 422283 448357 44629 453922 424046 43923 42927 411089 436511 428969 428969 428969 428969 428969 428969 428969 428969 428969 428969	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF053306 AF027866 AW238299 AI553633 NM_001898 AA456454 AA721252 AF120274 AF120274 AF120274 AF120274 T97307	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.214291 Hs.36708 Hs.3650618 Hs.326447 Hs.123114 Hs.214291 Hs.291502 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689	Human DNA sequence from clone RP5-850E9 NM_0210481Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (milotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1 serine (or cysteine) proteinase Inhibito UL16 binding protein 2 ESTs cystatin SN cell division cycle 2-like 1 (PITSLRE pr ESTs artemin artemin artemin artemin artemin artemin artemin phye53h05.s1 Soares fetal liver spleen hypothetical protein XP_098151 (leucine- NM_002362:Homo sapiens metanoma antigen,
556065707580	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 223 & 230 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 236 & 237 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 245 Seq ID No: 245 Seq ID No: 250 & 251 Seq ID No: 250 & 251 Seq ID No: 254 & 255 Seq ID No: 256 & 257 Seq ID No: 266 & 257 Seq ID No: 268 & 259 Seq ID No: 269 & 261 Seq ID No: 269 & 261 Seq ID No: 260 & 261 Seq ID No: 260 & 261 Seq ID No: 262 & 263 Seq ID No: 264 & 265 Seq ID No: 268 & 259 Seq ID No: 274 & 275 Seq ID No: 278 & 279 Seq ID No: 278 & 277 Seq ID No: 278 & 277 Seq ID No: 280 & 281 Seq ID No: 282 Seq ID No: 282 Seq ID No: 282 Seq ID No: 285 & 286	114346 438956 40446 103312 320843 429065 446102 330495 413573 428479 332180 437915 441553 331692 441553 441553 442924 422283 448357 44629 4539223 429248 439223 429248 439257 411089 428969	W00847 NM_005025 Y12642 BE069288 AI753247 AW168007 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF053306 AF027866 AW238299 AI553633 NM_001898 AA456454 AA721252 AF120274 AF120274 AF120274 AF120274 T97307 AA648459 H39860	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.201877 Hs.114311 Hs.108923 Hs.279682 Hs.36708 Hs.138202 Hs.250618 Hs.326447 Hs.123114 Hs.214291 Hs.291502 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689	Human DNA sequence from clone RP5-850E9 NM_0210481Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleotide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (mitotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1 serine (or cysteine) proteinase inhibito UL16 binding protein 2 ESTs cystatin SN cell division cycle 2-like 1 (PITSLRE pr ESTs artemin artemin artemin artemin artemin artemin artemin artemin hypothetical protein AF301222 hypothetical protein AF3012121 (leucine-
5560657075	Seq ID No: 221 & 222 Seq ID No: 223 & 224 Seq ID No: 225 & 226 Seq ID No: 227 & 228 Seq ID No: 227 & 228 Seq ID No: 231 & 232 Seq ID No: 231 & 232 Seq ID No: 234 & 235 Seq ID No: 238 Seq ID No: 238 Seq ID No: 238 Seq ID No: 241 & 242 Seq ID No: 241 & 242 Seq ID No: 243 & 244 Seq ID No: 245 Seq ID No: 250 & 251 Seq ID No: 252 & 253 Seq ID No: 254 & 255 Seq ID No: 256 & 257 Seq ID No: 266 & 267 Seq ID No: 268 & 269 Seq ID No: 270 & 271 Seq ID No: 276 & 277 Seq ID No: 278 & 279 Seq ID No: 278 & 279 Seq ID No: 282 & 281 Seq ID No: 283 & 284	114346 438956 40446 415669 103312 320843 42905 413573 428479 332180 437915 441553 331692 429413 422283 448357 44629 453922 424046 43923 42927 411089 436511 428969 428969 428969 428969 428969 428969 428969 428969 428969 428969	W00847 NM_005025 Y12642 BE069288 AI753247 AW168067 U47924 AI733859 Y00272 Y00272 AF134160 AI637993 AA281219 AI683487 NM_014058 AW411307 N20169 AF081497 AA236776 AF053306 AF027866 AW238299 AI553633 NM_001898 AA456454 AA721252 AF120274	Hs.135056 Hs.78589 Hs.3185 Hs.34744 Hs.29643 Hs.317694 Hs.71642 Hs.149089 Hs.334562 Hs.7327 Hs.202312 Hs.121296 Hs.152213 Hs.202313 Hs.201314 Hs.108923 Hs.152213 Hs.279662 Hs.736708 Hs.36708 Hs.36708 Hs.138202 Hs.250618 Hs.326447 Hs.123114 Hs.214291 Hs.291502 Hs.291608 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.194689 Hs.385951 Hs.288467	Human DNA sequence from clone RP5-850E9 NM_0210481Homo sapiens melanoma antigen, serine (or cysteine) proteinase inhibito lysosomal Homo sapiens mRNA; cDNA DKFZp547C136 (fr Homo sapiens cDNA FLJ13103 fis, clone NT ESTs guanine nucleolide binding protein (G pr ESTs cell division cycle 2, G1 to S and G2 to cell division cycle 2; G1 to S and G2 to claudin 1 Homo sapiens clone N11 NTera2D1 teratoca ESTs wingless-type MMTV integration site fami DESC1 protein CDC45 (cell division cycle 45, S.cerevis RAB38, member RAS oncogene family Rh type C glycoprotein MAD2 (milotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1 serine (or cysteine) proteinase Inhibito UL16 binding protein 2 ESTs cystatin SN cell division cycle 2-like 1 (PITSLRE pr ESTs artemin artemin artemin artemin artemin artemin artemin phye53h05.s1 Soares fetal liver spleen hypothetical protein XP_098151 (leucine- NM_002362:Homo sapiens metanoma antigen,

	W U 02/000				takan ata amataka Busan antakata anta
	Seq ID No: 293 & 294	424629	M90656	Hs.151393	glutamate-cysteine ligase, catalytic sub
	Seq ID No: 295 & 296	437789	Al581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
	Seq ID No: 297 & 298	437789	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheli
_	Seq ID No: 299 & 300	437789	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
5	Seq ID No: 301 & 302	437789	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheli
	Seq ID No: 303 & 304	437789	Al581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
	Seq ID No: 305 & 306	453968	AA847843	Hs.62711	High mobility group (nonhistone chromoso
	Seq ID No: 307 & 308	403478		40==00	NM_022342:Homo sapiens kinesin protein 9
10	Seq ID No: 309	441525	AW241867	Hs.127728	ESTs
10	Seq ID No: 310 & 311	434105	AW952124	Hs.13094	presentlins associated rhomboid-like pro
	Seq ID No: 312 & 313	428810	AF068236	Hs.193788	nitric oxide synthase 2A (inducible, hep
	Seq ID No: 314 & 315	413691	AB023173	Hs.75478	ATPase, Class VI, type 11B
	Seq ID No: 316 & 317	423934	U89995	Hs.159234	torkhead box E1 (thyroid transcription f
1.0	Seq ID No: 318 & 319	409228	R16811	Hs.22010	ESTs, Wealdy similar to 2109260A B cell
15	Seq ID No: 320 & 321	425734	AF056209	Hs.159396	peptidylglycine alpha-amidating monooxyg
•	Seq ID No: 322 & 323	413582	AW295647	Hs.71331	hypothetical protein MGC5350
	Seq ID No: 324 & 325	438403	AA806607	Hs.292206	ESTs
	Seq ID No: 326 & 327	403329			unnamed protein product [Homo sapiens]
20	Seq ID No: 328 & 329	409893	AW247090	Hs.57101	minichromosome maintenance deficient (S.
20	Seq ID No: 330 & 331	119073	BE245360	Hs.279477	v-ets erythroblastosis virus E26 oncogen
	Seq ID No: 332 & 333	113195	H83265	Hs.8881	ESTs, Wealty similar to S41044 chromosom
	Seq ID No: 334 & 335	102283	AW161552	Hs.83381	guanine nucleotide binding protein 11
	Seq ID No: 336 & 337	101345	NM_005795	Hs.152175	calcitonin receptor-like
0.5	Seq ID No: 338 & 339	103280	U84722	Hs.76206	cadherin 5, type 2, VE-cadherin (vascula
25	Seq ID No: 340 & 341	102012	BE259035	Hs.118400	singed (Drosophila)-like (sea urchin fas
	Seq ID No: 342 & 343	105729	H46612	Hs.293815	Homo sapiens HSPC285 mRNA, partial cds
	Seq ID No: 344 & 345	134299	AW580939	Hs.97199	complement component C1q receptor
	Seq ID No: 346 & 347	412719	AW016610	Hs.816	ESTs
	Seq ID No: 348 & 349	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL
30	Seq ID No: 350 & 351	128924	BE279383	Hs.26557	plakophilin 3
	Seq ID No: 352 & 353	100486	T19006	Hs.10842	RAN, member RAS oncogene family
	Seq ID No: 354 & 355	419121	AA374372	Hs.89626 -	parathyroid hormone-like hormone
	Seq ID No: 356 & 357	409459	D86407	Hs.54481	low density lipoprotein receptor-related
25	Seq ID No: 358 & 359	330493	M27826		endogenous retroviral protease
35	Seq ID No: 360 & 361	417866	AW067903	Hs.82772	collagen, type XI, alpha 1
	Seq ID No: 362 & 363	418113	Al272141	Hs.83484	SRY (sex determining region Y)-box 4
	Seq ID No: 364 & 365	437016	AU076916	Hs.5398	guanine monphosphate synthetase
	Seq ID No: 366 & 367	429612	AF062649	Hs.252587	pituitary tumor-transforming 1
40	Seq ID No: 368 & 369	440704	M69241	Hs.162	insulin-like growth factor binding prote
40	Seq ID No: 370 & 371	431221	AA449015	Hs.286145	SRB7 (suppressor of RNA polymerase B, ye
	Seq ID No: 372 & 373	431565	AF161470	Hs.260622	butyrate-induced transcript 1
	Seq ID No: 374 & 375	431565	AF161470	Hs.260622	butyrate-induced transcript 1
	Seq ID No: 376 & 377	132354	BE185289	Hs.1076	small proline-rich protein 1B (comifin)
15	Seq ID No: 378 & 379	424441	X14850	Hs.147097	H2A histone family, member X
45	Seq ID No: 380 & 381	103768	AF086009	Hs.296398	gb:Homo sapiens full length insert cDNA
	Seq ID No: 382 & 383	417512	X76534	Hs.82226	glycoprotein (transmembrane) nmb
-	Seq ID No: 384 & 385	425266	J00077	Hs.155421	alpha-fetoprotein
	Seq ID No: 386 & 387	424503	NM_002205	Hs.149609	integrin, alpha 5 (fibronectin receptor,
50	Seq ID No: 388 & 389	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin
50	Seq ID No: 390 & 391	418007	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial
	Seq ID No: 392 & 393	418007	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial
	Seq ID No: 394 & 395	418738	AW388633	Hs.6682	solute carrier family 7, (cationic amino tissue factor pathway inhibitor 2
	Seq ID No: 396 & 397	415138	C18356	Hs.295944	G protein-coupled receptor 39
55	Seq ID No: 398 & 399	418506	AA084248 D13666	Hs.85339	in the common terms of the
22	Seq ID No: 400 & 401	423961		Hs.136348	perfostin (OSF-2os) monokine induced by gamma interferon
	Seq ID No: 402 & 403 Seq ID No: 404 & 405	414812 417433	X72755 BE270266	Hs.77367 Hs.82128	5T4 oncofetal trophoblast glycoprotein
		417433	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein
	Seq ID No: 406 & 407 Seq ID No: 408 & 409	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse
60		428227	AA321649	Hs.2248	small inducible cytokine subfamily B (Cy
,00	Seq ID No: 410 & 411	444381	BE387335	Hs.283713	ESTs, Weakly similar to S64054 hypotheti
	Seq ID No: 412 & 413 Seq ID No: 414 & 415	400303	AA242758	Hs.79136	LIV-1 protein, estrogen regulated
	Seq ID No: 416 & 417	411789	AF245505	Hs.72157	Adlican
	Seq ID No: 418 & 419	428698	AA852773	Hs.334838	KIAA1866 protein
65 ·	Seq ID No: 420 & 421	450098	W27249	Hs.8109	hypothetical protein FLJ21080
05	Seq ID No: 422 & 423	421552	AF026692	Hs.105700	secreted frizzled-related protein 4
	Seq ID No: 424 & 425	452747	BE153855	Hs.61460	lg superfamily receptor LNIR
	Seq ID No: 426 & 427	450375	AA009647	12.01.00	a disintegrin and metalloproteinase doma
	Seq ID No: 428 & 429	426215	AW963419	Hs.155223	stanniocalcin 2
70	Seq ID No: 430 & 431	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin
. •	Seq ID No: 432 & 433	432201	AI538613	Hs.298241	Transmembrane protease, serine 3
	Seq ID No: 434 & 435	427585	D31152	Hs.179729	collagen, type X, alpha 1 (Schmid metaph
	Seq ID No: 436 & 437	442117	AW664964	Hs.128899	ESTs: hypothetical protein for IMAGE:447
	Seq ID No: 438 & 439	431211	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn
75	Seq ID No: 440 & 441	447033	Al357412	Hs.157601	ESTs
. •	Seq ID No: 442 & 443	447033	AJ357412	Hs.157601	EST8
	Seq ID No: 444 & 445	447033	Al357412	Hs.157601	ESTs
	Seq ID No: 446 & 447	115522	BE614387	Hs.333893	c-Myc target JPO1
	Seq ID No: 448 & 449	410418	D31382	Hs.63325	transmembrane protease, serine 4
80	Seq ID No: 450 & 451	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119
	Seq ID No: 452 & 453	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119
	Seq ID No: 454 & 455	452461	N78223	Hs.108106	transcription factor
	Seq ID No: 456 & 457	412420	AL035668	Hs.73853	bone morphogenetic protein 2
	Seq ID No: 458 & 459	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara
85	Seq ID No: 460 & 461	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon
	*				

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	WO 02/080	5443			
	Seq ID No: 462 & 463	437852	BE001838	Hs.256897	ESTs, Wealty similar to dJ365O12.1 [H.sa
	Seq ID No: 464 & 465	402075			ENSP00000251056*:Plasma membrane calcium
	Seq ID No: 466 & 467	421110	AJ250717	Hs.1355	cathepsin E
_	Seq ID No: 468 & 469	451668	Z43948	Hs.326444	cartilage acidic protein 1
5	Seq ID No: 470 & 471	451668	Z43948	Hs.326444	cartilage acidic protein 1
	Seq ID No: 472 & 473	451668	Z43948	Hs.326444	cartilage acidic protein 1
	Seq ID No: 474 & 475	422282	AF019225	Hs.114309	apolipoprotein L
	Seq ID No: 476 & 477	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily mamber
10	Seq ID No: 478 & 479	439738	BE246502	Hs.9598	sema domain, immunoglobulin domain (Ig),
10	Seq ID No: 480 & 481	427747	AW411425	Hs.180655	serine/threonine kinase 12
	Seq ID No: 482 & 483	420281	Al623693	Hs.323494	Predicted cation efflux pump C15000305:gij3806122 gb AAC69198.1 (AF0
	Seq ID No: 484 & 485	405932			C15000305:gij3806122jgbjAAC69198.1] (AF0
	Seq ID No: 486 & 487	405932	A184 044200	Hs.10887	similar to lysosome-associated membrane
15	Seq ID No: 488 & 489	444342	NM_014398	Hs.103982	small inducible cytokine subfamily B (Cy
13	Seq JD No: 490 & 491	421379	Y15221 U65590	Hs.81134	Interleukin 1 receptor antagonist
	Seq ID No: 492 & 493	417079 430890	X54232	Hs.2699	glypican 1
	Seq ID No: 494 & 495 Seq ID No: 496 & 497	419721	NM_001650	Hs.288650	aquaporin 4
	Seq ID No: 498 & 499	444471	AB020684	Hs.11217	KIAA0877 protein
20	Seq ID No: 500 & 501	413063	AL035737	Hs.75184	chitinase 3-like 1 (cartilage glycoprote
20	Seq ID No: 502 & 503	433800	AI034361	Hs.135150	lung type-I cell membrane-associated gly
	Seq ID No: 504 & 505	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro
	Seq ID No: 506 & 507	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro
	Seq ID No: 508 & 509	450001	NM_001044	Hs.406	solute carrier family 6 (neurotransmitte
25	Seq ID No: 510 & 511	410407	X66839	Hs.63287	carbonic anhydrase IX
	Seq ID No: 512 & 513	309931	AW341683		gb:hd13d01.x1 Soares_NFL_T_GBC_S1 Homo s
	Seq ID No: 514 & 515	412719	AW016610	Hs.816	ESTs
	Seg ID No: 516 & 517	417034	NM_006183	Hs.80962	neurotensin
	Seq ID No: 518 & 519	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam
30	Seq ID No: 520 & 521	413753	U17760	Hs.75517	laminin, beta 3 (nicein (125kD), kalinin
	Seq ID No: 522 & 523	425650	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen
	Seq ID No: 524 & 525	423673	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage
	Seq ID No: 526 & 527	418663	AK001100	Hs.41690	desmocollin 3
25	Seq ID No: 528 & 529	418663	AK001100	Hs.41690	desmocollin 3
35	Seq ID No: 530 & 531	429610	AB024937	Hs.211092	LUNX protein; PLUNC (palate lung and nas
	Seq ID No: 532 & 533	406690	M29540	Hs.220529	carcinoembryonic antigen-related cell ad
	Seq ID No: 534 & 535	431846	BE019924	Hs.271580	uroplakin 1B protease inhibitor 3, skin-derived (SKAL
	Seq ID No: 536 & 537	422158	L10343	Hs.112341	cadherin 3, type 1, P-cadherin (placenta
40	Seq ID No: 538 & 539	431958	X63629 AL035864	Hs.2877 Hs.69517	differentially expressed in Fanconi's an
40	Seq ID No: 540 & 541	437044 428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino
	Seq ID No: 542 & 543	429211	AF052693	Hs.198249	gap junction protein, beta 5 (connexin 3
	Seq ID No: 544 & 545 Seq ID No: 546 & 547	417389	BE260964	Hs.82045	midkine (neurite growth-promoting factor
	Seq ID No: 548 & 549	431009	BE149762	Hs.48956	gap Junction protein, beta 6 (connexin 3
45	Seq ID No: 550 & 551	417542	J04129	Hs.82269	progestagen-associated endometrial prote
73	Seq ID No: 552 & 553	449230	BE613348	Hs.211579	melanoma cell adhesion molecule
	Seq ID No: 554 & 555	410555	U92649	Hs.64311	a disintegrin and metalloproteinase doma
	Seq ID No: 556 & 557	410555	U92649	Hs.64311	a disintegrin and metalloproteinase doma
	Seq ID No: 558 & 559	424687	J05070	Hs.151738	matrix metalloproteinase 9 (gelatinase B
50	Seq ID No: 560 & 561	418462	BE001596	Hs.85266	integrin, beta 4
-	Seq ID No: 562 & 563	410274	AA381807	Hs.61762	hypoxia-inducible protein 2 ·
	Seq ID No: 564 & 565	439606	W79123	Hs.58561	G protein-coupled receptor 87
	Seq ID No: 566 & 567	404877			NM_005365:Homo sapiens melanoma antigen,
~ ~	Seq ID No: 568 & 569	444781	NM_014400	Hs.11950	GPI-anchored metastasis-associated prote
55	Seq ID No: 570 & 571	418543	NM_005329	Hs.85962	hyaluronan synthase 3
	Seq ID No: 572 & 573	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 574 & 575	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t protein tyrosine phosphatase, receptor-t
	Seq ID No: 576 & 577	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
60	Seq ID No: 578 & 579	415817	U88967	Hs.78867 Hs.78867	protein tyrosine phosphatase, receptor-t
00	Seq ID No: 580 & 581	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 582 & 583	415817 421817	U88967 AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR
	Seq ID No: 584 & 585 Seq ID No: 586 & 587	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
	Seq ID No: 588 & 589	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
65	Seq ID No: 590 & 591	409420		Hs.54451	laminin, gamma 2 (nicein (100kD), kalini
05	Seq ID No: 592 & 593	332180	AF134160	Hs.7327	claudin 1
	Seq ID No: 594 & 595	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,
	Seq ID No: 596 & 597	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,
	Seq ID No: 598 & 599	439223	AW238299	Hs.250618	UL16 binding protein 2
70	Seq ID No: 600 & 601	409757	NM_001898	Hs.123114	cystatin SN
	Seq ID No: 602 & 603	428969	AF120274	Hs.194689	artemin
	Seq ID No: 604 & 605	428969	AF120274	Hs.194689	artemin
	Seq ID No: 606 & 607	428969	AF120274	Hs.194689	artemin
	Seq ID No: 608 & 609	428969	AF120274	Hs.194689	artemin
75	Seq ID No: 610 & 611	450701	H39960	Hs.288467	hypothetical protein XP_098151 (leucine-
	Seq ID No: 612 & 613	450701	H39960	Hs.288467	hypothetical protein XP_098151 (leucine-
	Seq 1D No: 614 & 615	414774	X02419	Hs.77274	plasminogen activator, urokinase
	Seq ID No: 616 & 617	407944	R34008	Hs.239727	desmocollin 2
QΛ	Seq ID No: 618 & 619	407944	R34008	Hs.239727	desmocollin 2
80	Seq ID No: 620 & 621	457489	A1693815	Hs.127179 Hs.99376	cryptic gene ESTs
	Seq ID No: 622 & 623	429547	AW009166	1 13.3331 U	gb:Human nonspecific crossreacting antig
	Seq ID No: 624 & 625	407242	M18728 M18728		gb:Human nonspecific crossreacting antig
	Seq ID No: 626 & 627 Seq ID No: 628 & 629	407242 407242	M18728		gb:Human nonspecific crossreacting antig
0.5		444006	BE395085	Hs.10086	type I transmembrane protein Fn14
85	Seq ID No: 630 & 631				

WO 02/086443 PCT/US02/12476 Seq ID No: 632 & 633 NM_003816 Hs.2442 a disintegrin and metalloproteinase doma Seq ID No: 634 & 635 gastrin-releasing peptide 422109 S73265 Hs.1473 Seq ID No: 636 & 637 AW470411 Hs.288433 neurotrimin 419235 similar to S68401 (cattle) glucose induc Seq ID No: 638 & 639 Z45051 Hs.22920 449048 5 Seq ID No. 640 & 641 Hs.164021 small inducible cytokine subfamily B (Cy 419216 AU076718 Seq ID No: 642 & 643 Hs.256311 granin-like neuroendocrine peptide precu 431462 AW583672 Seq ID No: 644 & 645 448243 Hs.52620 AW369771 integrin, beta 8 Seg ID No: 646 & 647 426427 M86699 Hs.169840 TTK protein kinase Seq ID No: 648 & 649 445537 AJ245671 Hs.12844 EGF-like-domain, multiple 6 10 Seq ID No: 650 & 651 422278 AF072873 Hs.114218 frizzled (Drosophila) homolog 6 Seq ID No: 652 & 653 428450 NM_014791 Hs.184339 KIAA0175 gene product Seq ID No: 654 & 655 446619 AU076643 Hs.313 secreted phosphoprotein 1 (osteoponiin, Seq ID No: 656 & 657 453392 U23752 Hs.32964 SRY (sex determining region Y)-box 11 Seq ID No: 658 & 659 426514 BE616633 Hs.170195 bone morphogenetic protein 7 (osteogenic 15 Seq ID No: 660 & 661 425776 U25128 Hs.159499 parathyroid hormone receptor 2 Seq ID No: 662 & 663 425776 U25128 Hs.159499 parathyroid hormone receptor 2 Seq ID No: 664 & 665 431515 NM_012152 Hs.258583 endothelial differentiation, lysophospha Seq ID No: 666 & 667 419452 U33635 Hs.90572 PTK7 protein tyrosine kinase 7 Seq ID No: 668 & 669 432653 N62096 Hs.293185 ESTs, Weakly similar to JC7328 amino aci 20 Seq ID No: 670 & 671 432653 N62096 Hs.293185 ESTs, Weakly similar to JC7328 amino aci Seq ID No: 672 & 673 432653 N62096 Hs.293185 ESTs, Wealdy similar to JC7328 amino aci Seq ID No: 674 & 675 432653 N62096 Hs.293185 ESTs, Wealty similar to JC7328 amino aci Seq ID No: 676 & 677 410001 AB041036 Hs.57771 kallikrein 11 Seq ID No: 678 & 679 426501 AW043782 Hs.293616 **ESTs** 25 solute carrier family 15 (H??? transport Seq ID No: 680 & 681 408369 R38438 Hs.182575 Seq ID No: 682 & 683 445413 AA151342 Hs.12677 CGI-147 protein prostate differentiation factor Seq ID No: 684 & 685 422424 AI186431 Hs.296638 matrix metalloproteinase 7 (matrilysin, Seq ID No: 686 & 687 428330 L22524 Hs.2256 Seq ID No: 688 & 689 420610 AI683183 Hs.99348 distal-less homeo box 5 30 TABLE 15B Unique Eos probeset identifier number CAT number: Gene cluster number 35 Accession: Genbank accession numbers **CAT Number** Accession Pkey 309931 AW341683 M27826 R78416 AA307645 AW957879 AW957800 AA633529 H03662 330493 33264_5 AL133916 N79113 AF086101 N76721 AW950828 AA364013 AW955684 Al346341 Al867454 N54784 Al655270 Al421279 AW014882 40 439285 47065_1 ALT39916 N/9113 AF086101 N/6721 AW990828 AA364013 AW990828 AA36413 AW990821 AW67652 N62351 N69253 AA626243 AI341407 BE175639 AA456968 AI358918 AA457077
AA009847 AA131254 AA374293 AW954405 H04410 AW660284 AA161166 BE157467 BE157601 H04384 W46291 AW663674 H04021 H01532
AA190993 H03231 H59605 H01642 AA852876 AA113758 AA626915 AA746952 AI161014 AA099554 R69067
AW118072 AI631982 T15734 AA224195 AI701458 W20198 F26326 AA890570 N90552 AW071907 AI671352 AI375892 T03517 R88265 450375 83327_1 451320 86576_1 Al124088 AA224388 Al084316 Al354686 T33652 Al140719 Al720211 T03490 Al372637 T15415 AW205836 AA630384 T03515 T33230 45 AA017131 AA443303 T33623 AI222556 T33511 T33785 AI419606 D55612 TABLE 15C 50 Unique number corresponding to an Eos probeset Pkey: Sequence source. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA sequence of human chromosome 22." Dunham I. et al., Nature (1999) 402:489-495. Ref: Indicates DNA strand from which exons were predicted. Strand: 55 Indicates nucleotide positions of predicted exons. Nt_position: Strand Nt position Pkey 402075 8117407 121907-122035,122804-122921,124019-124161,124455-124610,125672-126076 Plus 403329 8516120 96450-96598 Plus 60 116458-116564 403478 9958258 Plus 404440 7528051 80430-81581 Plus 404877 1095-2107 1519284 Plus 405770 2735037 61057-62075 Plus

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			GGATCTACCT				360
15			ACCTACTGTT				420
			AGAAGGGGAT				480
	CCGCCCTGGC	CCCGGGTGTC	CCCAGCCTGC	GCGGGCCGCT	TCCAGTCCCC	GGTGGATATC	540
	CGCCCCCAGC	TCGCCGCCTT	CTGCCCGGCC	CTGCGCCCCC	TGGAACTCCT	GGGCTTCCAG	600
20			GCGCCTGCGC				660
20			TCTGGGTCCC				720
	CTGCACTGGG	GGGCTGCAGG	TCGTCCGGGC	TCGGAGCACA	CTGTGGAAGG	CCACCGTTTC	780
			TCACCTCAGC CGTGTTGGCC				840 900
			GTCTCGCTTG				960
25			ATCTGCACTC				1020
	TATGAGGGGT	CTCTGACTAC	ACCGCCCTGT	GCCCAGGGTG	TCATCTGGAC	TGTGTTTAAC	1080
	CAGACAGTGA	TGCTGAGTGC	TAAGCAGCTC	CACACCCTCT	CTGACACCCT	GIGGGGACCT	1140
	GGTGACTCTC	GGCTACAGCT	GAACTTCCGA	GCGACGCAGC	CTTTGAATGG	GCGAGTGATT	1200
20	GAGGCCTCCT	TCCCTGCTGG	AGTGGACAGC	AGTCCTCGGG	CTGCTGAGCC	AGTCCAGCTG	1260
30	AATTCCTGCC	TGGCTGCTGG	TGACATCCTA	GCCCTGGTTT	TTGGCCTCCT	TTTTGCTGTC	1320
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			EDLPGEEDLP				120
			WRYGGDPPWP				180
			SVQLTLPPGL				240
45	VEGHRFPAEI	HVVHLSTAFA	RVDEALGRPG	GLAVLAAFLE	EGPEENSAYE	QLLSRLEEIA	300
-10							
-15	EEGSETQVPG	LDISALLPSD	FSRYFQYEGS	LTTPPCAQGV	IWTVFNQTVM	LSAKQLHTLS	360
45	EEGSETOVPG DTLWGPGDSR	LDISALLPSD LQLNFRATQP	LNGRVIEASF	LTTPPCAQGV PAGVDSSPRA	IWTVFNQTVM	LSAKQLHTLS	360 420
	EEGSETOVPG DTLWGPGDSR	LDISALLPSD LQLNFRATQP		LTTPPCAQGV PAGVDSSPRA	IWTVFNQTVM	LSAKQLHTLS	
50	EEGSETQVPG DTLWGPGDSR GLLFAVTSVA	LDISALLPSD LQLNFRATQP	LNGRVIEASF RGTKGGVSYR	LTTPPCAQGV PAGVDSSPRA	IWTVFNQTVM	LSAKQLHTLS	
	EEGSETQVPG DTLWGPGDSR GLLFAVTSVA Seq ID NO:	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque	LNGRVIEASF RGTKGGVSYR	LTTPPCAQGV PAGVDSSPRA PAEVAETGA	IWTVFNQTVM	LSAKQLHTLS	
	EEGSETQVPG DTLWGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Act	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque	LNGRVIEASF RGTKGGVSYR ence #: BC01392	LTTPPCAQGV PAGVDSSPRA PAEVAETGA	IWTVFNQTVM	LSAKQLHTLS	
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50	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Act Coding sequ 1 AGCGGGGTTG	LDISALLPSD LQINFRATQP FLVQMRRQHR 3 DNA seque 1d Accession bence: 438-1	LNGRVIEASF RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG	IWTVFNQTVM ABPVQLNSCL 41 TTGTCAAGGC	LSAKQLHTLS AAGDILALVF 51 AGAGAAGAGA	
50	EEGSETQVPG DTLWGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession dence: 438-3	LNGRVIEASF RGTKGGVSYR ence 1 #: BC01392 1391	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA	IWTVFNQTVM ABPVQLNSCL 41 TTGTCAAGGC GACTAGGACT	LSAKQLHTLS AAGDILALVF 51 AGAGAAGAAGA GAGAAGAAGA	420
50 55	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Ac: Coding sequ 1 AGGGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession Dence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA	LTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCGGCCAGAG	IWTVFNQTVM AEPVQLNSCL 41 TTGTCAAGGC GACTAGGACT GAGGAGGGAA	LSAKQLHTLS AAGDILALVF 51 AGAGAAGAA GAGAGAAGA TTCCAAAAAA GCGCTTTTT	60 120 180 240
50	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession ence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA CTCTCTCTTT	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCGGCCAGAG TTTTCCCCCA	IWTVFNQTVM AEPVQLNSCL 41 TTGTCAAGGC GACTAGGACT GCTTAAGCCT GCTTAAGCCT AGAGGAGGAA AATTATTCTT	LSAKQLHTLS AAGDILALVF 51 AGAGAAGAGA GAGAAGAGA GAGAAAGA GTCCAAAAAA GCGCTTTTTT CGCCTGATTT	60 120 180 240 300
50 55	EEGSETQVPG DTIWGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGGTGTTCCTCGCGGA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession dence: 438-1 11 1 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC GCCTGCGCT	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCCGGCAGGT CTCTCTCTTTCTCTCTCTCTCTCTCTCTCTCTCTCT	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTCCCCAG CCGCCAGAGAG CCGCCCGCCT	IWTVFNQTVM ABPVQLNSCL 41 TTGTCAAGGC GACTAGGACT GCTTAAGCCT GAGGAGGAA AATTATTCTT CCCCTCCTCC	LSAKQLHTLS AAGDILALVF 51 AGAGAAGAA GAGAAAAAA GCGCTTTTTT CCCCCCG	60 120 180 240 300 360
50 55	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT TCCTCGCGGA CCCGCGGGCC	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession Dence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC GCCCTGCGCT CCCCAAAGTC	LNGRVIEASF RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCCGCACACCC CCGGCCGGGC	LTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCTGCCAGAG TTTTCCCCCA CCGCCGCCT CGAGGGTCGG	AT TOTCAAGGC GACTAGGACT GAGGAGGAA AATTATTCTT CCCCTCCC CGGCCGCGG	S1 GAGAGAAGAGAGAGAGAAAGAGAGAAAAAAGGCTTTTTTTCCCCCCGCCGGGCCCGGGCCCGGGCCGGCCGCCGGCCGCCGGCCGCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCCGGCCGGCCC	60 120 180 240 300 360 420
50 55 60	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGGGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT TCCTCGCGGA CCCGCGGCC CCGCGCACAG	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession ence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC CCCCAAAGTC CGCCCGCATG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA CTCTCTCTTT CCCGACACCC CCCGGCCGGGC TACAACATGA	LTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCGGCCAGAG TTTTCCCCCA CCGCCCGCCT CGAGGGTCGG TGGAGACGGA	A1 TTGTCAAGGC GACTAGGACTG GAGGAGGGAA AATTATTCTT CCCCTCCTCC CGGCCGCGG GCTGAAGCCG	S1 AGAGAAGAA GAGAGAAAGA TTCCAAAAAA GCGCTTTTT CGCCTGATTT TCTCCCCCC CGGGCCGGCCCCCCCCCC	60 120 180 240 300 360
50 55 60	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT TCCTCGCGGA CCCGCGGGCC CCGCGGGCACA AGCAAACTTC	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession Dence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC GCCCTGCGCT CGCCCCAAAGT GGGGGGGCGGC GGGGGGGGGG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCCGGGC TACAACATGA GGCGCAACCT	HTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCGGCCAGAG TTTTCCCCCA CCGCCGCCTCCGAGGGTCCG TGGAGACGGAC TGGAGACGGA CCACCGCGGC	A1 TTGTCAAGGC GACTAAGCCT GAGGAGGAA AATTATTCTT CCCCTCCTCC GGCGCCGCG GCTGAAGCCG GCGCCGCG	S1 GAGAGAAGA GAGAGAAAGA TTCCAAAAAA GCGCTTTTT CGCCTGATTT TCTCCCCCG CCGGCCCGC CCGGCCCGC CGGCACCAGA	60 120 180 240 360 420 480
50 55	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG AGAGGAGAGA TAATAATAAC TCATCCTGAT TCCTCGCGGA CCCGCGGACAG AGCAAACTTC AAAACAGCCC AAGCGCGCACAA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque Ld Accession Dence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC GCCCTGCGT CCCCAAAGTC CGCCGCATG GGGGGGGGGG	LNGRVIEASF RGTKGGVSYR ence 1 #: BC01392 L391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCCGCAGGC CCCGCCCGGCCAGGC TACAACATGA GGCGCAACCT AGAGGAACCT AGAGGGCCCCA GAGAACCCCA GAGAACCCCA	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTGCCAGAG TTTTCCCCCA CCGCCCCCCCCCC	41 TTGTCAAGGC GACTAGACT GAGAGGGAA AATTATTCT CCCCTCCC CGGCCGC GCTGAAGCCG GCTGAGCCG GCTGAGCCG GCTGAGCCC CATGGTGTGG CTCCGAGATC	S1 AGAGAAGAGA GAGAGAAGA TTCCAAAAAA GCGCTTTTT TCTCCCCCG CGGGCCGGC CGGGCCGGC CGGGCCGGC CGCAACCAGA TCCCGCGGGC AGCAAGCGCC	60 120 180 240 300 420 480 540
50 55 60	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGGGGGGTTG GTGTTTGCAA AGAGGAGAG TAATAATAAC TGATCCTGAT TCCTCGCGGA CCCGCGCACAG AGCAAACTTC AAAACAGCCC AGCGGCCCAA TGGGCGCCAA TGGGCGCCAA TGGGCGCCAA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession ence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC CCCCAAAGTC CGCCCGCATG GGGGGGGGG GGACCGCGTC GGTGGAAACTT	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GGCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCCGGGC TACAACATGA GGCGCAACT AAGCGCCCA GAGAACCCCA TTGTCGGAGA	LTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCTGCCAGGCCAG	A1 TTGTCAAGGC GACTAGGACT GAGGAGGGAA AATTATTCT CCCCTCCTC CGGCGCGG GCTGAAGCCG GCTGAAGCCG GCTGAAGCCG GCTGAAGCCG GCTGCAGCCGC CATGGTGTGG CTCGGAGTC GCCGTCATC	S1 AGAGAAGAG GAGAGAAGA TTCCAAAAAA GCGCTTTTT TCTCCCCCG CGGCCCGC GGCAACCAGA TCCCGCGGC GGCAACCAGA AGCAGCCC GACGAGCCC GACGAGCCC GACGAGCCC	60 120 180 240 300 420 480 540 600 720
50 55 60	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Ac: Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT TCCTCGCGGA ACCGGGCCC CCGCGCACAG AGCAAACTTC AAAACAGCCC AGCGGCGCGAAA AGGGGCGCGAA AGCGGCGCAA AGCGGCCGCAA AGCGGCCCCAA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession Dence: 438-1 11 TCTATTAACT AAGGGGGAAA GAAAGAAAGG AATCATCGGC TCCAGTTTGC GCCCTGCGCT CCCCAAAGT CGCCGCATG GGGGGGGGGG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCCGGGC TACAACATGA GGCGCAACT AAGCGGCCAA TAGAACGAACT ATGTCGGAGA ATGAAGAGGC	HTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAGA TCTTCCCCA CCGCCGCCT CGAGGGTCGG CGAGGGTCGG TGAAGACGGA CCACCGCGCT TGAATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA AGATGCACTA ACACCGGATTA	A1 TTGTCAAGGC GACTAGGACT GAGGAGGGAA AATTATTCTT CCCCTCCTCC GCGCGCGG GCTGAAGCCG GCTGAAGCCG GCTGAAGCCG GCGGCCGG CTGGAGTT CCCCTCTCC CTCCTCCTCC CTCCTCCTCC CTCCTC	S1 GAGAGAAGA TCCCCGGGGC AGCAGAGACACCACAAACAA CCCCGGGGC CCCGGCCCACACCACAACAACAACACCACAACACACAC	60 120 300 360 420 540 600 660 720 780
50556065	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAT TCCTCGCGGA CCCGCGCACAG CCGCGCACAG AGCAAACTTC AAACAGCCC AGCGGCGCAA TGGGCGCGAA AGCGGCGCGAA AGCGGCGCGAA AGCGGCGCGAA AACCAAGAC	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession dence: 438-1 11 1 TCTATTAACT AAGGGGAAA GAAAGAAGG AATCATCGGC GCCCGCATTGC CCCCAAAGTC CGCCGCATG GGGGGGCGGC GGACCGCGTC GATGGCCCAG GTGGAAACTA AGGGCTGCAG GTGGAAACTA GACGCTTCAG GCTCATGAAG GCTCATGAAG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT CCCGACACCC CCGGCCGGC TACAACATGA GCGGCACCA AGCGGCACCA AGGAACCCC AAGAACCCC AAGAACCCCA ATGTAGGAGAC AAGGAACCCA ATGTAGGAGAC AAGAACACACA AAGAACACACA AAGAACACACA AAGAACCCCA AAGAACACACA	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTCCCCAG CCGCCAGAGAGAGCGA CCACCAGCGC TGAAGACGGA CGACAGAGAAGCA ACACGAGTTA ACACGCTGCC ACACGAGTTA ACACGCTGCC	A1 TTGTCAAGGC GACTAGACCT GCTTAATCTT CCCCTCCTC CGGCCGG GCTGAAGCCG GCTGAAGCCG GCTGAGCGG CTGAGCGG CTGAGCGG CTGAGCGG CTGAGCGG CTGAGCCGC CATGGTGTGG CTCGGAGAT CCCCTCCTC CGGCGCCGG CTGGGGCGGC CATGGTGTGG CTCGGAGAT CCCGTTCAT TAAATACCGG CGGCGGCTG	S1 AGAGAAGAA AGAGAAGAA AGAGAAAAA CGCTTTTT TCTCCCCCG CGGGCCCGC CGGAACCAGA TCCGCGGGC AGCAACCAGA CCCCGGGGC AGCAACCAGA TCCGCGGGC AGCAACCAGA TCCGCGGGC CGCAACCAGA TCCCGCGGGC CGCACCCCG CGCCCCCC CGCCCCCC CGCCCCCC CGCCCCCC	60 120 180 360 420 480 660 720 840
50 55 60	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TCATCCTGAT TCCTCGCGGA CCGCGCACAG AGCAACTTC AAAACAGCC AAGCGCGCAA TGGGCGCGAA TGGGCGCGAA TGGGCGCGAA TGGGCGCGAAACCAACTAC AGCGGCGCAAACCAACTAC	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque d Accession hence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAGG AATCATCGGC TCCAGATTIG GCCCTGCGCT CCCCAAAGTC CGCACTGGGCT CGCACGGGCGGC GGACGGGCGGC GATGGCCCAG GTGGAAACTT AGCGCTGCAC GCTCATGAAG CCTCATGAAG CATGGCGAGC	LNGRVIEASF RGTKGGVSYR ence 1 #: BC01392 L391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCCGCCCGCCCGCCCGGCCAGCACACCC CCGGCCAGGC TACAACATGA GGCGCAACCT AGAGAACCCCA GAGAACCCCA ATGTCGGAGA ATGTCGGAGA ATGAGAGAGCT GGGGTCAGGG GGGGTCGGGG GGGGTCGGGG GGGGTCGGGG	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTGCCAGGGTCGC CGAGGGTCGC TGGAGACGAA TGAATGCACAA CGAGAGAGA CCACCGCGGT AGATGCACAA CGCAGAGAACGC ACCCGGATTA ACACGCTGCC TGGGGGTCGC	ATVFNQTVM AEPVQLNSCL AT TGTCAAGGC GCTTAAGCCT GAGGAGGAA AATTATTCTT CCCCTCCTCC CGGCCGCGG GCTGAAGCCG GCTGAAGCCG GCTGAAGCCG GCTGAGCTG CTCGGAGATC GCCGTTCATC TAAATACCGC CGCGGGCCTG CCTGGGGCCTC CCTGGGCGCCC CCTGGGCGCCC	S1 AGAGAAGAGA GAGGATTTT CCCCCCG CGGGCCGGC CGGGCCGGC CGCAACCAGA TCCCGCGGGC GGCAACCAGC AGCAACCAC AGCAACCAC AGCAACCAC CCCCGCGGGC CCCGCCGGC GCCACCCGCCGGC GCCACCCGCCGGC GCGAGCCCCG GCGGCCCCCG GCGCCCCCG	60 120 180 240 300 420 480 540 660 720 780 800
50556065	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TCATCCTGAT TCCTCGCGA ACCGGGGCC CCGCGACAG AGCAAACTTC AAAACAGCCC AAACAGCCGAAAC AGCGGCGCAAA AGCGGCTGCG AAACCAACGA AGCGGCTGCG AAACCAACGA AGCGCCAATAG AGCGCAATAG AGCGCAATAG	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque 1d Accession Dence: 438-1 11 TCTATTAACT AAGGGGAAA GAAAGAAAGG AATCATCGGC TCCCAAAGTC CGCCTGCGCT CGCCGCATG GGGGGGGGG GGACCGCGTC GATGGCCAG GTGGAAACTT AGGGCTGCAC CCTCATGAAG CCATCATGAAG CCATCATGAAG CCATGCGAGC CATGCGAGC CATGCGAGC CATGCGAGC CATGCGAGC CAGGTTACGCG CAGTTACGCG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT CCCGCCACACC TCCTCTTTT CCCGACACC TACAACATGA GGGGCACCT AAGCGGCCCA TTGTCGGAGA ATGAAGGAGC AAGGATCAGA AGGATAAGT GGGGTCGGGG CACATGAACC CACATGAACC	LTTPPCAQGV PAGVIDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCCAG TCTGCCCAGCGCT CGAGGGTCGG TGGAGACGAA CCACCGCGCT TGAATGCACAA CGCCGCCT TGAATGCACAA CGCCGCT TGAATGCACAA CGCAGGG TGAATGCACAA CGGAGAAGCG ACCCGGATTA ACACGCTGCC TGGGGCCCGG GCTGGAGCAA	ATVFNQTVM AEPVQLNSCL 41 TTGTCAAGGC GACTAGGACT GAGGAGGGAA AATTATTCT CCCCTCCCC CGGCCGCG GCTGAAGCCG GCTGAGCCG GCTGAGCCG GCTGAGCCG CATGGTGTG CTCGGAGATC GCCGTTCATC TAAATACCGG CGGCGGCGC CGGCGGCGC CCGGCGGCGC CCGGCGCGCC CCGGCAGCTAC	S1 AGAGAAGAGA GAGAGAAGAGA TTCCAAAAAA GCGCTTTTT TCTCCCCCG CGGGCCGGC CGGGCCGGC CGGGCCGGC GGCAACCAGA TCCCGCGGGC GGCAACCAGA TCCCGCGGGC GGCAACCAGA TCCCGCGGGC GGCAACCAGA CCCGGCGGA CCCGGCGGA CTGGCCCCCG GACGAGCTA CCCAGCGAGA CTGGCCCCCA AGCATGATCC	60 120 300 360 420 480 540 660 720 780 840 960
50556065	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ AGCGGGGTTG GTGTTTGCAA AGAGGAGAG TAATAATAAC TCCTCGCGGA CCCGCCACAG AGCAAACTTC AAACAGCCC AGCGGCCCA AGCGGCCGAA TGGGCGCCGA AGCGGCGCAAA TGGGCGCCGA AGCGACAGA AGCGGCTGCG AAACTAGAA AGCGCCTAGA AGCGCCAATAG AGCGCCAATAG AGCGCATGGA AGCGCATGGA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque d Accession dence: 438-3 11 1 1 TCTATTAACT AAGGGGAAA GAAAGAAAGG GATCATCAGG GCCCGCATG CCCCAAAGTC CCCCAAAGTC CGCCGCATG GGGGGGCGG GGACCGCGTC GATGGCCAG GTGGAACTT AGGGTTGAC GCTCATGAAG CATGGCAGC CAGTTACCCG GGGTTACCCG GGGGTACCCCG GGGCTACCCCG GGGGTTACCCCG GGGGTACCCCG GGGGTACCCCG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCCGGGC TACAACATGA GGGGCAACT AAGCGGCCCA ATGAACATGA ATGAAGAGA ATGAAGAGA ATGAAGAGA ATGAAGAGC CAAGAACC CAGCACCGG	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTTCCCCA CCGCCGCCT CGAGGGTCGG TCGAGGACGGA CCACCGCGCT TGAATGCCTT AGATGCACAA ACACGCGGC TGAATGCCTT AGATGCCTT AGATGCACAA ACACGCGGGC GCGGAGTAA ACACGCTGCC TGGGGCCAG GCTGGAGCAA GCCTCAATGC	41 TIGTCAAGGC GACTAGGACTA GACTAGACCT GAGGAGGGA AATTATTCTT CCCTCCTCC CGGCGCGG CCTGAAGCCG GCTGAAGCCG GCGGCGGCC CATGGTCTAC CCCGGCGCCGC CTCGGAGATC CCCGGCGCCGC CCCGCCCCCCCCCC	S1 AGAGAAGAA AGAGAAGAA AGAGCTTTTT TCTCCCCCG GGGCCGGC CCGGGGCCGGC GGCAACCAGA TCCCGCGGGC AGCAAGCACA ACCCGGGGCAACCAGA TCCCGCGGGC AGCAAGCACA ACCCGGGGAACCAGA CTGGCCCCG GGCGAACCAGA CTGGCCCCG GGCGAACCAGA	60 120 300 360 420 540 600 620 780 840 900 1020
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5055606570	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC CCGCGCGCACAGA TGGCGCGCACAGA TGGGGCGCAAACTC AAACAGCCC AGCGGCACAGA AGCGGCGCAAAC AGCGGCGCAAAC AGCGCCGAATAG AGCGCCAATAG AGCGCATGCA AGCGCATGCA AGCGCATGCA AGCACCATGCA TGGCCCTTCCTTCCTTCCTTCCTTCCTTCCTTCCTTCCT	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque d Accession dence: 438-3 11 1 1 1 TCTATTAACT AAGGGGAAA GAAAGAAGG GACAGCT CCCCAAAGTC CGCCGCATC GGGGGGGGGG	LNGRVIEASP RGTKGGVSYR ence 1 #: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GGCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCCGGC TACAACATGA GGCGCAACT AAGCGCCCA ATGAAGAGA ATGAAGAGC CACATGAACG CACATGAACG CACATGAACG CACATGAACG CACATGAACG CACATGAACG CACATGAACG CAGGTCGGC CACTACAGCA ACGGTGGTCA AGGGGCCCC ACCTACAGCA ACGGGGCCCC ACGTGGTCCA AGGGGGCCCC ACGTGGTCCA AGGGGCGCCCT	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTTCCCCA CCGCCGCCT CGAGGGTCGG TGAATGCCTA AGATGCATA ACACGCGGG TGAATGCCTT AGATGCATA ACACGCTGCC TGGGGCAGGA CGCGGATTA ACACGCTGCC TGGGGCAGA ACACGCTGCC TGGGCCAGA CCTCAATGC TGCAGTACAA TGCCTACTC TGCAGGCCGG GCTGGAGCAA TGCCTACTC TGCAGGCCGG CCCAGGCCCGG GCCAGGCCCGG	A1 TOTOTAMAGE ATTOTAMAGE GACTAGGACTA GACTAGGACTA GACTAGGACTA GACTAGGACTA AATTATTCTT CCCTCCTCC CGGCGCCGG CCTGAAGCCG CCTGAAGCCG CCTGAGCCGG CTCGAGGCTAC CCCGTCATC CCGGCGCTCATC CCGCGCCGC CCCGTCATC CCAGCAGCTAC CCAGCAGCTAC CCAGCAGCCAC CTCCATGACCC CAGCAGCGCC CAGCACCAGC CAGCACCCCCG CGCACCCCCC CGCACCCCCC CGCACCCCCC CGCACCCCCCC CGCACCCCCCC CGCACCCCCCC CGCACCCCCC CGCACCCCCCC CGCCCCCCCC	S1 AGAGAAGAA AGAGAAAGA TTCCAAAAAA GGCTTTTT TCTCCCCCG GGGGCCGGC CCGGGGCCAGA TCCCAGAGA TCCCAGAGA TCCCAGAGA TCCCGCGGGC GGCAACCAGA TCCCGCGGGC GGCAACCAGA TCCCGCGGGC GGCAACCAGA TCCCGCGGGA CCCCGGCGGA CCCCGGCGAA CCCCCGGCGAA ACCCCTGGCA ACCCCTGGCAGA ACCCCTGGCG GACATGATCA	60 120 300 360 420 480 540 600 720 780 840 900 1020 1080 1140 1260
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50556065707580	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC CCGGCGCACAGA TGGTCGGGA AGCAGACTC AAACAGCCC AGCGCCACAGA AGCGGCGCAAAA TGGGCGCGA AACCAAGC GCGGCAATAG AGCGCCATGCA AGCGCATGCA AGCAGCTTGG TTACCTGTT GCATGTATCT CCCAGCACTA TGACCACTT GCATGTATCT CCCAGCACTA TGACACTGT GGAAATGGG TCAAAAAAAA AACACAATC GGGAATGGA TCAAAAAAAA AACACAATC GGGAAGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAAGGGCC GGGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGAGGGCC GGGGAGGGCC GGGAGGGCC GGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGGAGGGCC GGGAGGGCC CCCACCACT CCCCACCACT CCCCACCACC CCCCACCACC CCCCACCACC CCCCACCA	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque dd Accession dence: 438-3 11 1 1 1 TCTATTAACT AAGGGGAAA GAAAGAAGG GACTGCGCT CCCCAAAGTC CGCCCGCATG GGGGGGGGG GGACCGCGTC GATGGCCCG GTGGAACTT AGGGTGCAC GCTCATGAAG CATTACGCG GGGCTACCAC CCCCACAGC CCCCACAGC CCCCACAGC CCCCCACAGC CCCCCCCC	LNGRVIEASP RGTKGGVSYR PROCE ##: BC01392 1391 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCGGCAGGA CTCTCTCTTT CCCGACACCC CCGGCAGGGC TACAACATGA GGGGCAACT AAGCGGCCA ATGAAGAGA ATGAAGAGC ATGAAGAGC CACATGAACG CACATGAACG CACATGAACG ATGAGGGCCCA CCTACAGC CACATGAACG ACGGGCCCT CACGTGCCCG ACGTGCCCG ACGGGCCCT AGGTGCCG AGGTGCCCA AGGGAACTG AGAGAACTG AGAGAACTG AGAGAACTG AGAGAACTG ACGGAACTG ACGGAACTG ACGGAACTG ACGGAACTG ACGGAACTG CCGGTGCCCC ACCTACAGC ACGGGCCCT CCGGTGCCCC ACGTGCCCC ACGGGCCCCT CCGGTGCCCC ACGGGCCCCT CCGGGGCCCT CCGGGGCCCCT CCGGGGCCCT CCGGGGCCCCT CCGGGGCCCCT CCGGGGCCCCT CCGGGGCCCCT CCGGGGCCCCT CCTTGTATAGGCCCTTGTATAGC	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTTCCCCA CCGCCGCCT CGAGGGTCGG TGAATGCCTT AGATGCATA ACACGCGGC TGAATGCCTT AGATGCATA ACACGCTGCC TGAGGAGAGCG ACCGGGATTA ACACGCTGCC TGAGGAGAGCG ACCGCGGATGA ACACGCGGCAGACCACACGCCACACACACACACACAC	41 TTGTCAAGGC GACTAGGACTA GACTAGGACGGA GATTAGCCT GAGGAGGGA AATTATTCTT CCCCTCCTCC CGGCCGG CCTGAAGCCG GCGGCGGC CATGGTCGGCGG CTCGAGACTG CCCGGCGCGC CGCAGCAGA CTCCATCATC CCAGCAGGC CAGCAGCAGA CTCCATCACC CAGCAGGAGATA CAGCACAGGAC CAGCAGAGAAA AATTATCAAA ATGACAGCACA ATGAGAAAAC AAGAAAACC AAGAAAACC AAGAAAACC AAGAAAACC AAGAAAACC AAGAAAACC AAGAAAACC AAGAAAACC	SAKQLHTLS AAGDILALVF 51 AGAGAAGAA GAGAAGAA TTCCAAAAAA CGCTTTTTT TCTCCCCCG CGGGCCCGC CCGGCGCCCGC GACAACAA TCCCAGAGA TCCCCCGGGC AGCAACCAGA TCCCGCGGC AGCAACCAGA TCCCGCGGC AGCAAGACCAC AGCATGATCA CCCCCGCGGA ACCCCTGCCA AGCATGATCA CCCCTGGCA ACCCCTGGCA CCCCTGTGC CAAAACCAG CCCCGTGG CAAAACCAG CCCGGTACCC TCGCAAAACCAG CCCGGTACCC TGCAAAACGAG CCCGGTACGC TGCAAAACGAG CCCGGTACGC TGCAAAACCAT TGGGGAGGTT CGAAAAACTT	60 120 180 240 300 420 480 540 660 720 780 960 1020 1140 1200 1320 1380 1440 1500 1560 1620
505560657075	EEGSETQVPG DTIMGPGDSR GLLFAVTSVA Seq ID NO: Nucleic Aci Coding sequ 1 AGCGGGGTTG GTGTTTGCAA AGAGGAGAGA TAATAATAAC TGATCCTGAT TCCTCGCGGA CCCGCGCACAG AGCAAACTTC AAACAGCCC AGCGCCATAGA AGCGCGCATAGA AGCGCCATGCA AGCGCCATGCA CCTACATGAA TGGCCCTTGC GCATGTATCT CCCAGCACTA CGCAAATGGA TCACAATGTA TGCTCTTGC GCATGTATCT CCCAGCACTA CACACATGTG TCACATGTA TCACACATGTG TCACACATCT CGCAGGAGTACCA CTTACATGTA TCACACATGTG TCACACATGTG TCACACATGTG TCACACATCT GGGAGGTCGC TTAAAAAAAA AACACCAATC GGGAGGGCG TTTAAAAGTT	LDISALLPSD LQLNFRATQP FLVQMRRQHR 3 DNA seque d Accession dence: 438-1 11	INGRVIEASF RGTKGGVSYR ence ##: BC01392 21 TGTTCAAAAA GTAGTTTGCT GAGAGAAGTT GCCGGCAGGC TCCTCTCTTT TCCGACACCC CCGGCCAGGC TACAACATGA GGCGGCACT AAGCGGCACCA AAGCACCCA AGGAACCCCA AGGACCCCA CTGTCGGGG CACATGAACG CAGCACCCG CAGCACCCG CAGCACCCG CAGCACCCG CAGCACCCG ACGAACTGAACG AGGACCCGG ACGACCCG ACGACCCG TTGTGTGGAGA TCGGTGCCCC ACGAACTGAACG TCGGTGCCCC TCGGTGCCCG TCAGGAACTGG AGAGGAACTGG AGAGGAGAGT TCACGCAAAA TKGGGGGACT TCCTTGTATAG GGTAGGAGCT CCTTGTATAG GGTAGGAGCT	LTTPPCAQGV PAGVDSSPRA PAEVAETGA 31 GTATCAGGAG GCCTCTTTAA TGAGCCCAG TCTTCCCCAG CCGCCAGAGGTCGG TGAAGACGAA CCACCACAGC TGAGGCCAAA GCCTCAATGC TGAGTACAA GCCTCAATGC TGCAGGCCGG CTGAGGCCGG CTGAGGCAGA CCCCCACAGC CACCCCGCCG CCACGCCGC CCACGCCGATGC AACCGCCGCAAA TGTCCTAATC AGTCCACAGC CCACACAGC CACCCACAGC ACCGCCGCATGC ATTGCAGGAA TTTCTGAGGA ATTGCAGGAAA TTTCTGAGGA TTTCCAGGAAA TTTCCAGGAAA	A1 TTGTCAAGGC GACTAGGACT GCTTAAGCCT GCTGAGGCGA GCTGAGCCG GCTGAGCCG GCTGAGCCG GCTGAGCCG CTGAGCCG CTGAGCCG CTGAGCCG CTGAGCCG CTGAGCCG CTGAGCCG CTCGGCCGC CTCGTCATC CCCTCCTC CCGCCGGCCGC CTCATGACC CCAGCAGCTAC CCACGCGCC CCCAGCAGA TAACCGG CCCCAGCAGA ATTTTCAAAA ATGGAGAAACTA AGAAAGCTA TTTGCAAAAG TTTTCAAAAG TTTTCCAAAAG TTTTCAAAAAGCTA TTTTCCAAAAGCTA TTTTCCAAAAAGCTA	SAKQLHTLS AAGDILALVF 51 AGAGAAGAA GAGAAAAAA GAGCTTTTT TCTCCCCCG CGGGCCCGC CGGGCCCGC CGCGCCCGC GGCAACCAGA TCCCGCGGC AGCAAGACC AGCATGATG TCCCCCTGTGG AGCATGATG CCCCCTGTGG ACCCCTCTGGCA CCCCCTTTGG CCCCTTTGG CCCCTTTGG CCCCTTTGG CCCCTTTGG CCCCTTTGG CCCGTACGCA CTTCACATGT CTGCCCCTCT GAAAAACCAG CCCGGTACGC CCCGGTACGC TGCAAAAACCAG CCCGGTACGC TGCAAAAACCAG TGGGGGGGGT TGGAAAAACTT TCTTTACCAA	60 120 300 360 420 540 600 600 780 960 1020 1080 1140 1260 1320 1440 1500 1560 1620 1620 1620 1620 1620
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WO 02/086443

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Seq ID NO: 59 Protein sequence:

WO 02/086443
Protein Accession #: NP_001784.2

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	PTSTATIVVH	VEDVNEARVF	VPPSKVVEVQ	EGIPTGEPVC	VYTAEDPDKE	NOKISYRILR	480
	DPAGWLAMDP	DSGQVTAVGT	LDREDEQFVR	NNIYEVMVLA	MDNGSPPTTG	TGTLLLTLID	540
1.5		RQITICNQSP					600
15		KQDTYDVHLS					660
		LVLLLLVRKK					720
		RNDVAPTIIP				APPYDTLLVF	780
	DYEGSGSDAA	SLSSLTSSAS	DQDQDYDYLN	EWGSRFKKLA	DMYGGGEDD		
20	Com TD NO.	60 DWA					
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	1	11	21	31	41	51	
25	ī	ī	ī	ī	ī	Ī	
	CCCTTCCCTT	GCCGCCGGAT	TCGAACGTTC	GGACTGAGGT	TTTTCTGCCT	GAAGAAGCGT	60
		GGATTGTTTT					120
		AAGCTGAGGC					180
	TCTCCCAGAG	GAAGCAGATA	AAGCGGAAGG	CTCCCCGTGG	CTTTCTAAAG	CGAGTCTTCA	240
30	AGCGAAAGAA	GCCTCAACTT	CGTCTGGAGA	AAAGTGGTGA	CTTATTGGTC	CATCTGAACT	300
		TGTTCATCGA					360
		TAACAAGGAG					420
	GAGGTTAGAA	GTCAAAGAAC	ATATTCTTGA	AAGTTATGAT	GCATTCTTTT	GGGTGGTAAC	480
25	AGATCATAAA	GACATTTTTT	ACACATCAGT	TAATATGGGA	TTATTAAATA	TTGG	
35							
		61 Protein			•		
	Protein Acc	cession #: F	os sequence	9			
40	1	11	21	31	41	51	
40		1					60
	-	KQIKRKAPRG		PQLKLEKSGD	PPAHPMCPPL	VARDARESKI	90
	NACASKCRVI	nkehvlaaak	VILKKSKG				
	Sea ID NO.	62 DNA goor	ence				
45		62 DNA sequ		194.2			
45	Nucleic Act	ld Accession	#: NW_0000	94.2			
45	Nucleic Act		#: NW_0000	94.2			
45	Nucleic Act	ld Accession	#: NW_0000	94.2 31	41	51	
	Nucleic Act	id Accession lence: 99-89	1 #: NM_0000		41 	51 	
45 50	Nucleic Act Coding sequent	id Accession Lence: 99-89	1 #: NM_0000 933 21]	31 	1	51 GCAAGGGACA <	
	Nucleic Acc Coding sequents 1 GGGCTGGAGG GAGGCGGGGG	id Accession nence: 99-89 11 GGCGCTGGGC TCCTAGCTGA	#: NM_0000 23 21 1 TCGGACCTGC CGGCTTTTAC	31 CAAGGCCACC TGCCTAGGAT	 GCAGGGGGGA GACGCTGCGG	GCAAGGGACA CTTCTGGTGG	120
	Nucleic Acc Coding sequence 1	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGGATC	#: NM_0000 233 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT	 GCAGGGGGGA GACGCTGCGG GCGAGCCCAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA	120 180
	Nucleic Actions sequently	id Accession Lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGGATC CACGCGCCTT	# : NM_0000 21 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT	 GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT	120 180 240
50	Nucleic According sequence of the control of the co	id Accession nence: 99-89 11 	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT	 GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT	120 180 240 300
	Nucleic According sequences of the control of the c	id Accession Lence: 99-89 11	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGGT TACGCCCCTG CGCAGGTCC CAGGGTGTGC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC	 GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC	120 180 240 300 360
50	Nucleic Actions sequently	id Accession nence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCCTT CAGCATTTC AGCCAGTGCA GTTCGGCCTG	#: NM_0000 333 21 1 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGGTGC CAGGGTGGC GATGCACTTG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG	 GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC	120 180 240 300 360 420
50	Nucleic Actions sequently	id Accession tence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCCTT CAGCAATTTC AGCCAGTGC GTTCGGCCTG CTACAAGGGG	#: NM_0000 21 TGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCTGC CAGGGTGTGC GATGCACTTG GGCAACACTC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCAC GCTCTGGGGG GCACAGGGGC	CAGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGAT TGCAATTCTC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG	120 180 240 300 360 420 480
50	Nucleic According sequence of the control of the co	id Accession tence: 99-89 11 	#: NM_0000 21 CGGACCTGC CGGCAGGG TACGCCGCTG CGCGAGGTCC CAGGGTGTGC CAGGACACTC CAGGCACCTC CAGGCCCCACCCC CAGGCCCCCC CAGGCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT GCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC	CAGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGAT TGCAATTCTC CAAGGTCTGC	GCAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATC CATGTGGCTG ATCCTGATCA	120 180 240 300 360 420 480 540
50	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGTGC GATGACACTC GGCAACACTC CTGGCCCGAC CTGGTGGACA	31 CAAGGCCACC TGCCTAGGAT CGCCCCAGGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCCC CAGCTGCCCA	CAGGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC CAAGGTCTGC AAGGCTGAAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATC CATGTGGCTG ATCCTGATCA GGGCAGGGG	120 180 240 300 360 420 480 540
50	Nucleic According sequence of the control of the co	id Accession tence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCCTT CAGCAATTTC AGCAATTTC CTACAAGGGC CTACAAGGGG CCTGCCCCAG GTCCCAGGAC TGCTGTGGGGG	#: NM_0000 333 21 1 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGC GATGCACTTG GGCAACACTC CTGGCCCGAC CTGGTCGAC ATCAAGAATG	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCCA CTGACCCTGA	CAGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGATGTGATC TGCAATTCTC CAAGGTCTGA GAGGCTGAA GGAGCTGAAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGG GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGG CGAGTTGCCT	120 180 240 300 360 420 480 540 600 660
50	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCATTTC AGCCAGTGC GTTCGGCCTG CTACAAGGGG CCTGCCCAG GTCCCAGGAC TGCTGTGGGG CTCCCAGGAC TGCTGTGGGG	#: NM_0000 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CAGGCACACTC CTGGCACACTC CTGGCCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCACTTGCCAC GCTCTGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA CTGACCTGA TCAATGACTT	CAGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTGA GGTGCAGTAC TGCAATTCTC CAAGGTCTGAC AAGGCTGAAG GGAGCTGAAG CAGCATCTTG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GGGCAGCGG CGAGTTGCCT AGCGAGCAGCGA ATCCTGATCA AGGACAGCAGC CAGGTTGCCT AGGACACCTAC	120 180 240 300 360 420 480 540
50	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGCGCTT CAGCAGTTCA GTTCGCCTG CTACAAGGGG CTCCAAGGGG CTCCCAGGAC TGCTGCCCAG GTCCCAGGAC TGCTGGGGC TTCCCGGACTT TTCCCGGAGAGAC	#: NM_0000 21 21 TCGGACCTGC CCGGCTTTTAC CTGGCAGGG TACGCCGCTG CGCGAGGTCC CAGGGTGTGC CAGGGTGTCC CAGGGTGTCC CAGGGTGTCC CAGGGTGTCC CAGGGTGTCC CTGGCCCGAC CTGGTCGACA ATCAAGAATG TTCTTCTCG GTGTGCACGA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCATTGCCAC GCTCTGGGGG CTAGGGGGC CTGGTGTCCC CAGCTGCCCA TCAACCCTGA TCAATGACTT CTGCTGGTGG	CAGGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGAATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG GCAGCTGAAG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG	GCAAGGACA CTTCTGGTGG GCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTT ATCTGCTG ATCTGCTG ATCTGATCA GGGCAGGG CGAGTTGCCT AGGACACTAC ACCCGACTTC	120 180 240 300 360 420 480 540 600 660 720
505560	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGGATC CACGCCTT CAGCAATTTC AGCAATTTC AGCAGTGCA GTTCGGCCTG CTACAAGGG CCTGCCCAG GTCCCAGGAC TGCTGTGGG CTCCGAGTT TCCCGAGTA GACCTTCCGGAGA GACCTTCCGGAGA GACCTTCCCGAGAC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCGAC TCTGGCCGAC TCTGTGGACA ATCAAGAATG TTCTTCTTCT CTGGTGGACA CCACGAGACCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCAGGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG CCACAGGGGC CTGGTGTCCC CAGCTGCCCA TCAACACTG TCAATGACT TGGTGGTG TGGTGTGTGTGTGTGTGTGTGTGTGTGT	CAGGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGAATTCTC CAAGGTCTGC AAGGTCTGC AAGGTCTGAAG CAGCATCTGC CAGGACTGAAG CAGCATCTTG TGAGCCCAAGG TGAAGCCAAGG	GCAAGGACA CTTCTGGTGG GCACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ATCCTGATCA AGCGACGTC ATCCTGATCA AGCGACCTC AGCGACCTC AGCCACCTC AGCCACCTC AGCCACTCC AGCCACTCC AGCCACTCC AGCCAATCCT	120 180 240 300 360 420 480 540 600 660 720 780
50	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCTT CAGCAATTTC CATCAGCTGCA GTTCGGCCTG CTACAAGGG CCTGCCCAG GTCCCAGGAC TGCTGTGGGG CTCCGACTTC TGCGGAGA CTCCGGAGA GTCCCGGAGA GCCCTGGCCGGGGACACTCC TGCGCGAGA	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGCC CAGGGTGCC CTGGCCGAC ATCAAGAATG TTCTTCTTCG GTCACCAC GTCACCACCACCACCACCACCACCACCACCACCACCACCAC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG CCAGCTGCCCC AGCTGCCCCA CTGACCCTGA TCAATGACTT CTGCTGGTGC CTGTGTGTCCC CTGTGTGTCCC CTGACCCTGA TCAATGACTT CTGCTGTGC CTGTGTGTCC CTGTGTGTCCC CTGTGACTGTC	CAGGGGGGA GACGCTGCGG GACGCTGCGG CTTACTGGAT CGAAGGCTGA TGATGTGATC TGATGTGATC TGAAGTCTC CAAGGTCTGC AAGGCTGAA GGAGCTGAA GGAGCTGAA CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ACCCTGATCA AGGGAGGG CGAGTTGCCT AGGACACTAC ACCCGACCTC CAGGACATCC CAGGACATCC CAGGACATCC CAGGACATCC CAGGACATCCT CAGTACACTC	120 180 240 300 360 420 480 540 660 720 780 840
50556065	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGCGCTT CAGCAATTTC AGCAATTTC AGCAGTGCA GTTCGGCCTA GTTCGGCCTA GTCCCCAG GTCCCAGGC CTCCCAGGC CTCCGACTT CTCCGGAGA GACCTCTGCT GTGGACAGC GCTGGGACA GTTGGGCCCAG	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGGG TACGCCGCTG CGCAGGTCC CAGGGTGTCC CAGGCCGAC CTGGCCCGAC CTGGCCCGAC CTGGTGGACA TCAAGAATG TTCTTCTCG GTGTGCACGA CCACGAGACC CCACGAGACC CCACGAGACC CCACGAGACC CCACGAGACC CCACGAGACC CCACTGCCCA CCACTGCCCA CCACTGCCCAC CCACTGCCAC CCACTGCCCAC CCACTGCCAC CCACTGCCCAC CCACTGCCAC CCACTGCCAC CCACTGCCAC CCACTGCCCAC CCACTGCCAC CCACTGCCCCAC CCACTGCCCCAC CCACTCCCAC CCACTCCCCAC CCACTCCCAC CCACTCCCAC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA TCAATGACTT CTGCTGGTGG TCGTTGGTGG TGGTGTGCC GTGACCTGA GGTGACTGC GTGACCTGA	CAGGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC CGAGGTCTGC AAGGTCTGC AAGGCTGAAG GCAGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC CGAGGTCAAG CGAGGTCAAC CGAGGTCAAC CGAGGTCAAC CGAGGTCAAC CGAGTACCAA	GCAAGGACA CTTCTGGTGG GCACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GGGCATCC AGGACACTAC AGCCACTC AGGACACTAC ACCCGACCTC AGCCAATCCT CAGTACACTC GTCCCAGCTG GTGACTCG	120 180 240 300 360 420 540 600 660 720 780 840 900
50556065	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGCGCTT CAGCAATTTC AGCAATTTC AGCAGTGCA GTTCGGCCTG GTCCCCAG GTCCCAGGC CTCCAGGAC TTCCGCGTGC GTCCGACTT GTGGGCG GTCGGACTT GTGGACAGC GCTGGGACAG GCTGGGACAG GCTGGGACAG GCCAACAGC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGGG TACGCCGCTG CGCAGGTCC CAGGGTGTCC CAGGGTGTCC CTGGCCCGAC CTGGTCGACA TCAAGAATG TTCTTCTCG GTGTGCACGA CCACGAGACC CCACGAGACC CACGAGACC CACGAGACC CCACGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGAGACC ATCAGGGAGACC ATCAGGGAGACC ATCAGGGAGACC ATCAGGGAGACC ATCAGGGAGACC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA TCAATGACTT CTGCTGGTGG TCGTTGGTGG TCGTGGTGG TCGTGGTGG TCGTGACCTGA GCCACTGACCTGAC	CAGGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGAATGTGATC CAAGGTCTGC AAGGCTGAAG GCAGCTGAAG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCATCATG CAGCAAGC CTACAAGGTC CGAGGTCAAG GAAGCTCAAG GAAGCTCAAG GACAGCTCAG	GCAAGGACA CTTCTGGTGG GCACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGCATGCC CATGTGGCTG ATCCTGATCA GGGCAGGGG CGAGTTGCCT AGGACACTAC ACCCGACCTC AGCCAATCCT CAGTACACTC GTCCCAGCTG GTCCCAGCTG ATCCTGATCA ACCCGACCTC AGCCATCCT AGCCAATCCT AGCCAATCCT AGCCAATCCT AGCCCAGCTC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020
50556065	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC CTCCTAGCTGA CGCGGATC CACGGCCTT CAGCAATTTC CAGCAATTTC CTACAGGG CTTCCAGGAC GTCCCAGGA CTCCAGGAC TGCTGGGG CTCCGACTTC TTCCCGGAGA GACCTCTGCT GTGGACAGC GCTGGCCAG GCTGGCCAG GCTGGCACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GGGACAGG GGAACTGACC	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGTCC CAGGGTGCAC ATCAAGAATG TTCTTCTTCG GTGACAC CTGTGCACA CCACGAGAC CCACGAGAC CCACTGCCGA CCACTGCCGA CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGTCTCC CACGGGACA ATCCAGGATA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCTGAT TCAATGACTT CTGCTGGTGG TGGTGCTGC TGGTGCTGGC GGCACAGGCCC GGCACAGGCCC CTGTGACTGG CTGAGCGCC CTGTGACCGC CTGTGACCGC	CAGGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGATGTGATC CAAGGTCTGC AAGGCTGAA GAGCTGAAG GAGCTGAAG CAGCATCTTG CTGCCTGTG TGAGCCAAGC CTACAAGGTC CGAGGTGAAC CAAGCTCACG CAACACCCAGC CAACCCCCGG CAGCCTCCTG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GGGCAGGGG GGAGTTGCCT AGGACACTAC ACCCGACCTC AGCCAATCCT AGCCAATCCT GGCCAATCCT GGCCAATCCT GGCCAGCTG GTGACTGTGA ACCACTGCCC GTGGCCTGGCC	120 180 240 300 360 420 480 540 600 660 720 780 840 900 960 1020 1080 1140
50556065	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGATTC CACGGGATTC CACGGGATTC AGCCAGTGCA GTTCCAGGAC GTCCCAGGA GTCCCAGGAC TCCTGCTCCGACTTC TCCCGACTTC TCCCGACTTC GTGGACAGC GCCCAACAGC GCTGCCCACACGC TGTGCCCACACACC GGGACTACC AGGTGCCACTC AGGTGCCACCTC AGGTGCACCTC AGGTGCACCC AGGTGCCACCTC AGGTGCACCC AGGTGCCACCTC AGGTGCCACCTC AGGTGCCACCTC AGGTGCCACCTC AGGTGCCACCTC AGGTGCACCC AGGTGCCACCTC AGGTGCCACCTC AGGTGCACCTC AGGTGCACCTC AGGTGCACCC AGGTGCACCC AGGTGCACCC AGGTGCACCC AGGTGCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTACCC AGGTGCCACCC AGGTCCACCC AGGTGCCACCC AGGTCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACCC AGGTGCCACC AGGTGCCACCC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGAGTCTCC CTGGCAGAC CTGGCAGAC TTCTTCTTCG GTGTGACA TTCTTCTTCG GTGTGCACGA CCACGAGACC CCACGAGACC CCACGAGAC CCACGAGAC CCACTGCCGA CGCAGTGCCC ATCGGGGAGG ATCGGGGAGG ATCAGAATA GGCTACCGTG GGCATACCGTG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGGCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC AGCTGCCA TCAATGACTT CTGCTGGTG TGGTGTGTG TGGTGTGTG TGGTGTGTG TGGTGACTG GTGACTGAC CTGTGACTG CTGTGACTG CTGTGACTG CTGTGACTG CTGTGACTG CTGTGACTG CTGACGCCC TGACAGCCCC TGACAGCCCC TGACAGCCCC TGACAGCCCC TGACAGCCCC TGACAGCCCC TGACATGGCG	CAGGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCTGAG AGTGCAGTAC TGATGTGATC CAAGGTCTGC AAGGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGGCCAAGC CTACAAGGTCAAC GCAGGTGAAC GAAGCACTCAG CAAGCTCAG GAAGCTCAG GAAGCTCAG GAAGCTCAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGAAGCAC CGAGTTGCCT AGGACACCTC CAGCACCTC CAGTACACTCC GTCCAGCTG GTGACTGCAGCTG GTGACTGCAGCTG GTGACTGCAGCTG GTGACTGCAGCTG GTGACTGCAC GTGACTGCAC GTGGCCCCA	120 180 240 300 360 420 480 540 600 660 720 780 900 960 1020 1080 1140 1200
50556065	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GCCGCTGGGC TCCTAGCTGA CCCGGGATC CACGCGCTT CAGCATTTC AGCCATTGC CTACAAGGGG CCTGCCCAG GTCCCAGGAC TGCTGCAGAC TGCTGCAGAC TGCTGGGCTT TCCCGGAGAC GCCGACTTC TGGGACAGC GCTGGCCAG GCTGGCCAG GCCGACTGC GCGACTGC GGAACTGACC GGAACTGACC GGAACTGACC GGAGCTGGCC GGAGCTGGCC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGAGTCC CAGGGTGTGC CTGGCCGAC CTGGCCGAC CTGGTGACAC TTCTTCTCG GTGTGCACGA TCAAGAATG CTCACGAGACC CCACTGCCGA CCCACTGCCGA CCCCGGGCTCCC CCCGGGGCGCC CCCGGGGGCGCC CCCGGGGGCGCC CCCGGGGGCGCC CCCGGGGGCGCC CCCGGGGGG	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTCT GCTTTGCCAC CTGACCTCA TCAACGGGC CTGACCTCA TCAACGCTG TCGTGTGCCA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGAC	GCAGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGAATTCTC CAAGGTCTGAA GGAGCTGAAG GGAGCTGAAG GGAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGGCCAAGC CTACAAGGTC CGAGGTGAAC CAACACTCCAG GAAGCTCAAG GAAGCTCAAG GAAGCTCAAG GACGCTCCAG GACGCTCCAG GGTCCTCAGT GCTCCCAGT GCTGCGTGAC	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GGGCAGGGG GCGAGTTCC CAGGTGACC CAGGTGCTG ATCCTGATCA ACCCGACCTC AGGCAATCCT AGGCAATCCT CAGTACACTC GTCCCAGCTG GTCCAGCTG GTGACTGCCC GTGGCCTGC GTGGCCCCC TTGGACCCTG	120 180 240 300 360 420 540 600 660 720 780 840 900 1020 1080 1140 1200 1260
50556065	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGCGCTT CAGCAATTTC AGCCAGTGCA GTTCGGCCTG CTACAAGGGG CTCCAAGGGG CTCCAAGGGC TTCCGCCAG GTCCCAGGAC TGCTGTGGG GTCCGACTTC GTGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GGAGTGCACT GGAGCTGGCT GGAGCTGGCC TGAGGTGACCC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGGG TACGCCGCTG CGCGAGGTCC CAGGAGGTCC CAGGAGGTCC CAGGAGGTCC CTGGCCCGAC CTGGCCCGAC CTGGTGGACA ATCAAGAAT CTCTTCTTCG GTGTGCACGA CCAGGAGACC CACGAGACC CACGAGACC CCAGTGGCC ATCGGGAGAC CCAGTGGCC CCAGTGGCC CCAGTGCCC CCAGTGGCC CCAGTGCCC CCAGTGCCC CCAGTACCAGAATA CGCTACCGGGAGG CCTGCGGGAGG CCTGCGGCAGG CTGAGCACCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG CTGGTGTCCC CAGCTGCCCA TCAACCCTGA TCAATGACTT CTGCTGGTGG TGGTGCTGC TGGTGCTGC TGGTGCTGC TGGTGCTGC CTGTGACCTGA CCTGTGACCTGC TGGTGCTGC TGGTGCGCA TCAACCCCA TGACATGAC CTGTGACCGC CTGTGACCGC CTGTGACCGC CTGTGACCGC CTGTGACCGC CTGTGACCGC CTGTGACCGC TGACATGCC TGACATGCC TTATTTGCCC	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGAATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG CAGCATCATG CAGCATCATG TGAGCCAAGC CTACAAGGTC TGAGCCAAGC CTACAAGGTC CGAGTACCAA GACAGCTCCTG GGTCCTCAG GGTCCTCAG GCTCCTCG CAGCTCCAG CAGCTCCAGC CAGTGTGGGC CAGTGTGGGG CAGCTCGGGC CAGTGTGGGG	GCAAGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCTT AGCGATGACC CGGGCCATCC CATGTGGCTG ATCCTGATCA AGCACACTC AGGACACTC AGCACTC AGCAATCCT AGCAATCCT AGCAATCCT CAGTACACTC AGCCAATCCT CAGTACACTC GTCCCAGCTG GTGACTGCCC GTGGCCTG CTGACTGCCC GTGGCCTGC CGTGGGCCCA CTTGGACCTC CGTGGCCTGC CGTGGGCCCACTT	120 180 240 300 360 420 540 600 660 720 840 900 900 1020 1140 1200 1260 1320
50556065	Nucleic According sequence of the control of the co	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCAATTTC CACGAGCTGC GTCCAGGA GTCCAGGG GTCCCAGGA TGCTGGGC TGCTGGGG GTCCCAGGA GTCCGACTTC TGCGGAGA GACTCTGGC GGGACACAGC GGGACTGC AGGTGCCACT GGAGCTGGC TGGGACTGC TGCGACTGC TGGGACTGC TGGGACTGC TGGACTGAC TGGAGCTGGC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC TGGGGTGGCC	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGC GATGCACTTG GGCAACACTC CTGGCCGAC ATCAAGAATG TTCTTCTTCG GTGACAC CCACTGCCGA CCACTGCCGA ATCAAGAATA GCCACTGCCGA CTGGGCAGG ATCCAGAATA GCCACTGCCGA CTGGGCAGG CCTTCTGTTG GTGGACAC CCTGGCCGG CCTTCTGTTG GTGACACC CCTTCTGTTG	31 CAAGGCCACC TGCCTAGGAT CGCCCCAGGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGC CTGGTGCCCA TGACCCTGA TCAATGACTT CTGCTGGTGG GTGGCGCA GTGACCGCA GCCACTGCCA TGACCGCCA TGACCGCCA TGACCGCCA TGACCGCCA TGACCCCA TGACCCCA TGACCCCA TGACCCCA TGACAGCCCA TGACAGCCCA TGACATGCCC TTATTTGGCCG AGCAGACCCCT AGCAGACCCCT AGCAGACCCCT AGCAGACCCCT	GCAGGGGGGA GACGCTGCGG GCAGCCCAG CTTACTGGAT CGAAGGCTGA TGATGTGATC TGATTCTC CAAGGTCTGC AAGGCTGAG GAGCTGAAG GCAGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG CTACAAGGTC CGAGGTGAAC CAAGCTCAGC CAACCCCGGTCCCTG GCTCCTCG GCTCCTCGG GCTCCTCGG CCAGCTCCAGT CCAGCTGGGG CCACCCGGTC CCACCCGGTC CCACCCGGTC CCACCCGGTC CCCCCGTC CCCCCGGTC CCCCCGTC CCCCCGGTC CCCCCGGTC CCCCCGGTC CCCCCGGTC CCCCCGGTC CCCCCGGTC CCCCCGGTC CCCCCGCTC CCCCCGGTC CCCCCGCTC CCCCGCTC CCCCCGCTC CCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCCGCT CCCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCGCTC CCCCCCCC	GCAAGGGACA CTTCTGGTGG CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ACCACTGGCATCC CAGGACACCT AGGACACTAC ACCCGACCTC CAGTACACTC GTCCAGCTG GTGACTGTGAC GTGACTGTGA TCCTGACTG CTGACTGCC GTGGCCTG CTGGCCCACTT ATCCTGGGCC ATCCTGGGCCCACTT ATCCTGGGCC	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020 1140 1200 1140 1260 1320
50556065	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCAATTTC AGCAATTTC AGCAGTGCA GTCCAGAG GTCCCAGGG CTCCAGGGC TGCCCAG GTCCCAGGG GTCCCAGGG TCCGACTTC TCCCGGAGA GACCTCTGC GTGGGCCAG GCTGGGACAG GCTGGGACAG GGGACTGGC GGAGTGCC GGAGTGGC TGAGGTGACC TGAGGTGACC TGAGGTGACC TGAGGTGACC TCGCACTGAC CCCCCTTTCC CCCCCTTTCC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGAGGTCC CAGGGTGTGC CTGGCCGAC ATCAGAATG TCTTCTTCT GCAGAGGC CCACTGCCGA CCACTGCCGA ATCAGAGATG CCACTGCCGA CGCAGTGCCG CCACTGCCGA CGCAGTGCCG CCACTGCCGA CGCGGAGG CCACTGCCGA CGCGGAGG CCACTGCCGA CGGGGTCTC CCACTGCCGA CGGGAGG CCACTGCCGA CGGGAGG CCACTGCCGA CGGGAGG CCACTGCCGA CGGGAGG CCTGGGCAGG CTGAGCACT CCTGGGCAGG CTGAGCACCT CCTGGCCAG CTTCTTTTG TGGAACTTCG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGGCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTG TGATGACTG GTGACTGC GTGACTGC GTGACTGC GTGACTGC GTGACTGC GTGACTGC GTGACTGC GTGACTGC GTCACAGCCCA TGACATGCCT TGCTGACTG TGTTAGTGT TATTTGGCCG TTCAGTGTT TATTTGGCCC TGCTGAGGC TGCCTGAGGC TGCCTGAGCC TGCCTGAGCC TGCCTGAGCC TGCCTGAGC TGCCTGAGCC TGCCTGAGC T	CAGGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCCTA AGTCAGTAC TGATGTGAT TGCAATTCTC CAAGGTCTGC AAGGCTGAA GGAGCTGAAG CAGCATCTTG CGTGCCTGTG CTACAAGGTCAG CAAGCTCAG GCAGCTGAAG CAAGCTCAG CAAGCTCAG CAAGCTCAG CAGCTCCTG GGTCCTCAGT CAGCTCCTG GGTCCTCAGT CCTGCGTGAC CAGTGTGCGGCCCCGGTC CCGTGGCTAC	GCAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGAGGAGCA GGACAGCGC CAGGACCAC ACCCGACCTC CAGCACCTC GTGCACTCC GTGCACTCC GTGCACTCC GTGCACTCC GTGCCATCCT GTGCACTCC GTGCCATCCT GTGCCACTG GTGGCCTGAC TTGGAGCCTG CCGCCACTT TGGAGCCTG CCGCCACTT TTGGAGCCTG CCGCCACTT ATCCTGGGCC CCGGTTGGAAT	120 180 240 300 360 420 480 540 600 660 720 960 1020 1140 1200 1140 1200 1380 1380 1440
5055606570	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GCGCTGGGC TCCTAGCTGA CGCGGATT CAGCATTTC AGCAATTTC AGCAATTTC AGCAGAC CTACAAGGG CTTACAAGGG CTGCCCAG ATCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCGGACTT CTGCGCCTG GTGGACAGG GCTGGGACAGG CGCAACAGC GGAACTGACC AGGGCTAGC GGAACTGACC AGGGCTAGC TCGCACTGC TCGCACTTC CGCACTTC CGCACTTC CGCACTTC CGCACTTC CGCACTTC CGCACTTC CCCCCTTTCC GACTGGCTTG CCCCCTCTTCC CGACTGCCTTGCC CGACTGGCTTGCC CCCCCTTTCC CGACTGGCTTGCC CGACTGGCTTTCC CGACTGCCTTTCC CGACTGGCTTTCC CGACTGGCTTTGCC CGACTGCTTTGCC CGACTGGCTTTGCC CGACTGGCTTTGCC CGACTGGCTTTGCC CGACTGGCTTGCC CGACTGGCTTGCC CGCACTGCTTTGCC CGACTGGCTTGCC CGACTGGCTTGCC CGCACTGCTTTGCC CGACTGGCTTGCC CGCACTGCTTTGCC CGCACTGCTTTGCC CGACTGCTTTGCC CGACTGGCTTTGCC CGACTGCTTTGCC CGACTGCTTTCC CGACTGCTTTCC CGACTGCTTTGCC CGACTGCTTTCC CGCACTGCTTTCC CGACTGCTTTCC CGACTGCTTTCC CGCACTGCTTTCC CGACTGCTTTCC CGACTGCTTTCC CGCACTGCTTTCC CGACTTTCC CGCACTGCTTTCC CGCACTGCTTTCC CGCACTTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTTCC CGCACTTCTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTCC CGCACTTTCC CGCACTTTCC CGCACTTTCC CGCACTTCC CGCACTTTCC CGCAC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CAGGAGACACTC CTGGCAGAC CTGGTGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA TCCGGGAGGC CCACTGCCGA TCCGGGAGG CCTCGCTCC CTTCTGTTC CTGGCAGC CCTTCTGTTC TGGAACTTGG GAGCCACCGC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTTGCCAC CTGACCTTGACCTGA CTGACCTTGA TCAATGACTT CTGCTGGTG TGGTGCTGACTGAC TGGACCTGAC TGACCTGAC TGACAGCCCA GGCACTGAC CTGTGACTGC GTGACTGCC GTTAACTGC GTTAACTGC GTTAACTGC GTTAACTGC GTTAACTGC AGCAGCCCA TGACAGCCCA TGACATGCCG TGACTGCCG TGACTGCCG TGTTAGTGTT TATTTGGCCG AGCAGACCCA AGCAGACCCA AGCAGACCCA AGCAGACCGA AGCAGACCGA AGCAGGCGCA AGCAGACCGCA AGCAGACCCCA AGCAGACCGCA AGCAGACCCA AGCAGACCGCA AGCAGACCCA AGCAGACCGCA AGCAGACCGCA AGCAGACCGA AGCAGACCGA AGCAGACGCA AGCAGACCA AGCAGACA AGCAGACCA AGCAGACCA AGCAGACCA AGCAGACCA AGCAGACA AGCAGACCA AGCAGACA AGCAGACCA AGCAGACA AGCAGACA AGCAGACA AGCAGACA AGCAGACCA AGCAGACA AGCAGACA AGCAGACA AGCAGACA AGCAGACA AGCAGACA AGCAGACCA AGCAGACA	GCAGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCTGAG CTGATTCTC CAAGGTCTGC AAGGCTGAG GGAGCTGAG GGAGCTGAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC CTACAAGGTC CTACAAGGTC CTACAAGGTC CGAGTACCAA GACACTCGG CAGCTCCGG GCGCCTCTGG GCTGCTTGGGGG CGCCCCGTTG CCGTGGGTGAC CAGTGTGGGG CCCCGTGGCTAC ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTC ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCTCTCT ACTGCCCTCTCT	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GCGCATCC CATGTGGCTG ATCCTGATCA GCGCAGCCTC AGCCAGCCTC AGCCAATCCT AGCCAATCCT GTGCCAGCTG GTGCCAGCTG GTGCCAGCTG GTGACTGCCC GTGGCCTGGC GTGGCCTGGC GGGGGCCCA TTGGAGCCTG ATCCTGGCCTGG	120 180 240 300 360 420 540 600 660 720 780 840 900 1020 1080 1140 1260 1320 1320 1340 1440 1500
50556065	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGGCCTT CAGCAATTC AGCAATTC GTTCGGCCTG GTTCGGCCTG GTCCCAGGAC TGCTGGGCCTG GTCCGAGTT TCCCGAGTT TCCCGAGTC TGCGCCAG GCTGGGCAGC GGACTGC GGACTGC GGACTGC GGACTGC TGCGCCACAC TGTGGGCTG TGCGCCAC TGCGCCTG TGCGCCTG TGCGCCTGC TGCGCCTGC TGCGCCTGC TGCGCCTGC TGCGCCTGCC TCGCACTGC TGAGTGCCC TCGCACTGC TGAGTGCCCTTG GGATGCCCTTGC GGATGCCTTG GGATGGCCTG	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCG CAGGGTGCC GATGCACTTG GGCAACACTC CTGGCCGAC ATCAAGAATG TCTTCTTCTC GTGTGCACGA CCACGAGGCC CCACTGCGA ATCAGAGAC GCCAGTGCCGA ATCAGAGAC GCCAGTGCCGA ATCAGAGAC GCCAGTGCCGA ATCAGAGAC GCCAGTGCCG GCTTCTCTTG GGGAACTTCCGTG GTGAGCACCC GCTTCTGTTG TGGAACTTCG CAGCCAGC CAGCCGCGACC CAGCCGGCAC CAGCCGCGCACC CAGCCGCGCACC CAGCCGCGCAC CAGCCGCGCCACCCC CAGCCGCGCCACCCC CAGCCGCGCCACCCC CAGCCGCGCCACCCC CAGCCGCGCCACCCCC CAGCCGCGCCACCCC CAGCCGCGCCACCCCC CAGCCGCGCCACCCC CAGCCGCCACCCC CAGCCGCGCCACCCC CAGCCGCCCCC CAGCCGCCCCC CAGCCGCCCCC CAGCCGCCCCCC CAGCCGCCCCC CAGCCGCCCCC CAGCCGCCCCC CAGCCGCCCCC CAGCCCGCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG CTGGTGTCCC CAGCTGCCCA TCAACCCTGA TCAATGACTT CTGCTGGTGC GTGAGCCGA TCAATGACTT CTGTGACCTGC GTGAGCGGC GTGAGCGGC CTGAGCGGC CTGAGCGCCA TGACATGAC CTGTGACCGG CTGAGCCCA TGACATGAC CTGTGACCGG CTACAGCCCA TGACATGGC TTATTTGGCC AGCAGACCCT TGCCTGAGG AGCAGACCCT TGCCTGAGGG AGCAGACCCT TGCCTGAGGG AGCAGACCCT TGCCTGAGGG CTGAGTACCG	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGAATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG CAGCATCATG CAGCTCAGA CAGCATCATG CGAGCTCAGA GACACCCCAG GACGTCCTCG GGAGGTCCTCAGT GCTCCTCAGT GCTCCTCAGT GCTCCTCAGT CCCTCACTCCTC CCTCACCTCCTC CCTCACCTCCTC CCTCACCTCCTC CCTCACCTC	GCAAGGACA CTTCTGGTGG GCACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGGGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGG CGAGTTGCCT AGCACACTC AGCCAATCC GTCCCAGCTG GTCCCAGCTG GTGACTGCC GTGGCCTGC GTGGCCTGA CCACTGCCC GTGGCCTGC CGTGGCCCA TTGGAGCCTG CCGCCACTT ATCCTGGGCC CGGTTGGAC CCGCCACTT ATCCTGGCC CGGTTGGAC CCGCCACTT ATCCTGGCC CGGTTGGAC CGGTTGGAC CCGCCACTT ATCCTGGCC CGTTGCACC CTACACTCTCC	120 180 240 300 360 420 540 600 660 720 780 840 900 1020 1080 1140 1260 1320 1380 1440 1500 1500
5055606570	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCTT CAGCAATTTC CACAGTGCA GTTCGGCCTG CTACAAGGG CCTGCCCAG GTCCCAGGA TTCTGGCCTG TGCTGGGC TGCTGGACATTC TGCGGACA GTCCGACACAC GTGGCACACAC GGAACTGAC GGAACTGAC CGCAACAG GGAACTGAC GGAGCTGGC TGGGCTGGC TGGGGTGGC TGGGACTGAC CGAGTGGCC GGAGTGGCC	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGC CAGGGTGC CTGGCCGAC ATCAAGAATG TCTTCTTCG GTGACAC ATCAAGAATG TCTTCTTCG GCCAGGCC CACTGCCGA ATCAGGACC CCACTGCCGA CGCAGGGCC CACTGCCGA ATCAGAATA GCTACCGTG CCTGGCAGG ATCCAGAATA GCTACCGTG CCTGGCAGC CCTTCTGTTG TGGAACTTCG CAGCCCCCCCAAACCCCC CACCCCCCAAACCCCCC CACCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCAGT GCAGCTTTCT GCATTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTGG TGGTGCTGT TGGTGGTGC CTGTGACCTGA GCCACTGAC CTGTGACCTGA CTGTGACCGA CTGAGCGCC CTGTGACCGA CTGTGACCGA CTGTGACCGA CTGTGACCGA CTGTGACCGA CTGTGACCGA CTGTGACCGA CTGTGACCCA TGACTGTT TATTTGGCCG AGCAGACCCT TGCCTGAGGC CTGGAGACCCT TGCCTGAGGC CCGTGGTTCC CCGTGGTTCC	GCAGGGGGGA GACGCTGCGG GCAGCCCAG CTTACTGGAT CGAAGGCTGA CGAAGGCTG TGATTCTC CAAGGTCTGC TGCAATTCTC CAAGGTCTGAG GCAGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG CTACAAGGT CCAAGGTCAAG CCAGCTCCTG GGTCCTCAGT GCTCCTCAGT GCTCCTCTG GCTCCTCAGT CCAGCTCCCCC CCACGGTCCCCCCCCCC	GCAAGGGACA CTTCTGGTGG CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ACCACTGGCCT AGGACACTCC CAGGACGACCT AGGACACTCC CAGTACACT GTGCAGTGGC GTGCCATCC GTCCAGCTG GTGACTGCC GTGGCCTG GTGGCCTG CCGGCCCACTT ATCCTGGAC CCGGTTGGAAT CTCTGGAC CTGGTTGAAT CTCTGGAC CTGGTTGAAT CTCTGGAC CTGGTTGAAT CTCTGGCC CGGTTGGAAT CTACACTCTCC CAGTTGCACT CAGTTGCAC CTTCTGGAC CTTCTGGAC CTTCTGGAC CTTCTTGAAC CTTCTTGAAC CTTCTTGGAC CTTCTTGAAC CTTCTTGGAC CGGTTTGAAT CTCTTGGCC CGGTTTGAAT CACTCTTCC CAGCTTCCC CGGTTGCACT CACTCTCCC CGGTTGCCTG CACCTCTCC CGGCTGCCTG CACCTCTCC CGGCTGCCTG CACCTCTCC CGGTTGCACT CTCTCTCC CGGCTGCCTG CTTCTCTCC CGGCTGCCTG CTTCTCTCC CACCTCTCC CACCTCTC CACCTCTCC CACCTCTCC CACCTCTC CACCTCTCC CACCTCTC CACCTCTCC CACCTCTCC CACCTCTCC CACCTCTCC CACCTCTCC CACCTCTC CACCTCTCC CACCTCTC CACCTCTCC CACCTCTC CACCTCT CACCTCTC CACCTCT CACCT CACCTCT CACCTCT CACCT CACC	120 180 240 300 360 420 540 660 720 840 900 900 1020 1140 1200 1260 1320 1380 1440 1500 1560 1620
5055606570	Nucleic According sequences of the control of the c	Id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA TCCTAGCTGA TCCAGGGATTC CACGGGATTC CACGGCCTT CAGCAATTTC CACCAGTGCA GTTCCAGGA GTCCCAGGA TCCCAGGA TCCCAGGACTTC TCCCGGAGA GACCTCTGCT TCCCGGACA GCTGGCCACACAG GCTGCCAACAG GCTGCCAACAG GCAACAGC GCAACAGC TGAGCTGGCC CACCACTGC CACCACTCC CACCACTCC CACCACTCC CACCACTCC CACCACTCC CACCACTCC CACCACTCC CACCACTCC CACCACC CACCACC CACCACC CACCACC CACCAC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC ATCAGAATA TCTTCTTCT GGCACACAC CCACTGCCGA CCACTGCGCAC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACCGCCACCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTG TGATGACT TGTGACCTGA TCAATGACT TGTGACCGC TGGACTGCCA GCCACAGCCCA GGCACTGAC TGACTGCC TGACTGCC TGACTGCC TGACTGCC TGACTGCC TGACTGCC TGACATGCC TGACATGCC TGCTGAGGC AGCAGCCC TGCCTGAGGC AGCAGCCC TGCCTGAGGC CCGTGGTTCC CCGTGGTTCC AGCTGCCCG	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCTGAG AGTGCAGTAC TGATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG GGAGCTGAAG CAGCATCTTG CGAGCTGAAG CAGCATCTTG CGAGCTGAAG CAGCATCTTG GGAGCTGAAG CAAGCTCGAG CAAGCTCAGC CGAGTGCAC GCAGCTCCTG GGTCCTCAGT GCTCCTGG GCTCCTCAGT CCGTGGCTAC CCGTGGCTAC CCGTGGCTAC CCTCACACTC CCTCACACTC CACTGGACCA GCAGCGGTG CACTGGACCA GCAGCGGTG CACTGGACCA GCAGCGGTC CCTCACACTC CACTGGACCA GCAGCGGTC CACTGGACCA GCAGCGGTC CCACTGGACCA GCAGCGGGTC CCACTGGACCA GCAGCGGTC CCACTGGACCA GCAGCGGCTC CCACTGGACCA GCAGCGGTC CACTGGACCA GCAGCGGTC CACTGGACCA GCAGCGGCTC CCACTGGACCA GCAGCGGCTC CACTGGACCA GCAGCGGTC CACTGGACCA GCAGCGGCTC CCACTGGACCA GCAGCGCGCC GCACGGCTC CACTGGACCA GCAGCGGCTC CACTGGACCA GCAGCGGCTC CACTGGACCA GCAGCGCGCC GCACGGCCC GCACGGCCC GCACGGCCC GCACGCGCC GCACCAC GCACC	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATTCC CATGTGGCTG ATCCTGATCA ACCAGCAGCAG GGAGAGGAG GGAGTGCCT AGGACACTAC ACCCGACCTC CAGTACACTCC GTGCCATCTC GTGCCAGCTG GTGACTGTGA ACCACTGCCC GTGGCCTG CCGCACTT ATCCTGGGCCC CGGTTGGAAT GATGTGAAT GATGTGAAC GAGTGCCCG CGGTTGGAAT CCGGCACTT CCGGCCCC CGGTTGGAAT GATGTGACCCC CGGTTGCACC CGGTTGCCTG CGAGTGCCTG CGAGTGCCTG	120 180 240 300 360 420 480 540 660 720 780 840 900 900 1020 1140 1200 1320 1380 1440 1560 1560 1620 1680
5055606570	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCATTTCA AGCCAGTGC CTACAAGGGG CTTACAAGGGG CTCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCGGACTT CTCCGGACTT CTCCGGACT GTGGCCACAGAC GCCAACAGC GGAACTGACC AGGTGCCAC GGAGCTGGCC TCGCACTGC TCGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CCCACTGC CCCACTGC CCCACTGC CCCACTGC CCCCACTGC CCCCACTGC CCCCCTGCC CCCCCCC CCCCCCC CCCCCCC CCCCCCCC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGAGTGTCC CAGGAGTGTCC CTGGCAGAC CTGGTGACA ATCAGAATG TTCTTCTTCG GTGTGCACGA CCACGAGACC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAATA GGCTACCGT CCTGGCAGA CCACTGCCGA ACCAGTACC CCTTCTTTG CGGCAGG CGCTTCTTTTG CAGCACCG CACCCGCCA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCCCTGCAA CCACAGAACC CCACTGCCGC ACCCCTGCAA CCACAGTACC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAACCCTGA TCAATGACTT CTGTGGTG GTGACCTGA GGCACAGCCCA GGCACAGCCCA GGCACAGCCCA TGACATGAC TGACTGAC GTTAGTGAC GTTAGTGAC GTTAGTGAC GTTAGTGAC GTTAGTGAC GTTCAGTGT TATTTGGCCG AGCAGACCC AGCAGCCCC AGCAGCCCC AGCAGCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC AGCAGCCCC CCGTGGTTCC CCGTGGTTCC CCGTGGTCCCGG GCATCATTGT	GCAGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTCAAA GACACTCAG GCAGCTCAG GCAGCTCAG GCTGCTGG GCTGCTGGG GCTCCTCAGT GCTGCGTGAC CAGTGTGGGG CAGCCTCTCT CCTCACACTC CCTCACACTC CACTGGACCA GCAGCGGGTGAC CACTGGACCA ACTGCCCTCT CCTCACACTC CACTGGACCAC GCAGCGGGTGAC GCAGCGGGTG GCGCAGCACCAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GCGCACCTC CATGTGGCTG ATCCTGATCA ACCCGACCTC AGCCAATCCT AGCCAATCCT AGCCAATCCT AGCCAATCCT GTGCCAGCTG GTGACTGCCC GTGGCCTGGC GTGGCCTGGC GTGGCCTGGC GTGGCCTG ATCCTGGCCT ATCCTGGCC CGGTTGGAAT GATGTGAAT GATGTGACCC TACACTCTCG CGGTTGGATTGAAT GATGTGACCC CGGTTGGATTGAAT GATGTGACCC CGGTTGGATTGAAT GATGTGACCC CACCTCTCC CAGCTGCTC CAGGTGCTCT CAGGGGGTTG CAGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTG CACGGGGGTTC CACGGGGTTC CACGGGGGTTC CACGGGGTTC CACGGGGTTC CACGGGGTTC CACGGGGTTC CACGGGGGTTC CACGGGGTTC CACGGGGTTC CACGGGGGTTC CACGGGGTTC CACGGGGTT CACGGGGTTC CACGGCACGT CACGGGGTTC CACGGGC	120 180 240 300 360 420 540 600 660 720 780 840 900 1020 1120 1260 1320 1320 1320 1440 1500 1680 1680 1740
505560657075	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATC CACGGCCTT CAGCAGTTC AGCAGTTCA GTTCGGCCTG GTCCCAGGA TGCTGCCCAG GTCCCAGGAC TGCTGGACAT TGCGACTTC GACTTC GACTTC GACTTC GACTTC GACTTC GGACTGC GGACTGC GGACTGC GGACTGCC TGCGCCACAC AGGTGCCACT GGACTGCC TGGACAC TGCGCCTT CCCACTGC CGACTGC CCCACTGC CCCACTGC CCCACTGC CCCCTTTCC GACTGCC GACTGGCC GACTGGCC GACTGGCC GCACTGGC CCCCTTTCC GACTGGCC CGCTGTCC CGACTGCC CCCTGGTCC CCCTGGTCC CCTGGTCC CCTGGTCC CCTGGTCC CCTGGTCC CCTGTTCC CGTGCTCC CCTCTTCC CGTGCTCC CGTGCTCC CGTGCTCC CGTGCTCC CGTGCTCC CGTGCTTCCT	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCG GATGCACTTG GGCAACACTC CTGGCCGAC ATCAAGAATG TCTTCTTCTC GTGTCACGA ATCAAGAATG TCTTCTTCTCC GTGTGCACGA CCACGAGGCC CCACTGCGA ATCAGAATA GGCTACCGT GTGACCGG ATCAGAATA GGCTACCGT GTGACCGG CTTCTGTTG TGGAACTTCG CAGCGGGCA ACCCCTGCAA ACCCCTGCAA CAAGCCACCG GCAGCACCC GCGGGCA ACCCCTGCAA CAAGCCACCG GCGGGCAA CAAGCCACCG GCGGGCAA CCAGGACACG	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT CGCTTTGCCAC GCTTTGCCAC GCTCTGGGGG CTGGTGTCCC CAGCTGCCCA TGACCCTGA TCAATGACTT CTGCTGGTGC TGGTGCCCA TGACCTGAC TGACCTGA TCAATGACTT CTGTGACTGC GTGACCTGA GCCAATGAC CTGTGACCGG GTCAGCCCA GCCACAGCCA AGCAGCCA AGCAGCCA AGCAGACCCT TGCCTGAGGC TGCCTGAGGC CCATGAGC CCATGAGCCCA AGCAGACCCT TGCCTGAGGC CCATGAGCCCA AGCAGACCCT CCGTGGTTCC AGCATCCCC CCGTGGTTCC CAGCATCCAC CCATCCACCG CCATCATTGT CAGCATTCGA	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCCTGC CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG GGAGCTGAAG GGAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGGCCAAGC CAGCTCCTG GCAGCTCCTG GCTGCTCGTG CGTGCTCAGT GCTCCTCAGT CCTCAGTGCCCTCTC CCTGGACCACCC CACTGGACCA CACTGGACCA CACTGGACCA CACTGGACCA CACTGGACCA CACTGGACCA CACTGGACCA CACTGGACCA CCTTGGACCA CCTTGGATGAC CCTTTGGATGAC CCTTACAGCCC CTTTGGATGAC CCTTACAGCCC CTTTGGATGAC CCTTACAGCC CCTTTGGATGAC CCTTACAGCCC CTTTGGATGAC CCTTACAGCCC CTTTGGATGAC CCTTACAGCCC CTTTGGATGAC CCTTACAGCACC CCTTTGGATGAC CCTTACAGCACC CTTTGGATGAC CTTTGGATGAC CCTTACAGCACC CTTTGGATGAC CTTACAGCACC CTTTGGATGAC CTTACAGCACC CTTTGGATGAC CTTACAGCACC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGATGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGATGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGATGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGGATGAC CTTTGATGAC CTTTTGATGAC CTTTGATGAC CTTTTGATGAC CTTTTTTTTTT	GCAAGGACA CTTCTGGTGG GCACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GGGCAGGGG GGAGTTGCCT AGCACACTC CAGTTGCTG ATCCTGATCA ACCCGACCTC AGCCAATCC GTGCCAGCTG GTGACTGCC GTGGCTGGCTG GTGACTGCC GTGGCCTGC CTGGCCTGC CTGGCCTGC CTGGCCTGC CTGGCCTGC CTGGCCTGC CCGCCACTT ATCCTGGGCC CGGTTGGAAT GATGTGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGCCTG CAGGGGTTG CCAGGGGTTG CCAGGGGTTG CTACACTCCTC CAGGGGGTTG CTTCAGGCTG CTTCAGCTG CTTCAGC	120 180 240 300 360 420 480 5540 600 660 720 780 840 900 1020 1140 1200 1320 1380 1440 1500 1560 1620 1680 1740 1800
5055606570	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCTT CAGCATTTC CAGCATTTC CTACAAGGG CTTCCAGAC GTCCCAGG GTCCCAGG GTCCCAGG CTCCCAGGC TGCCCAGG GTCCCAGGC TGCCCAGGC TGCCCAGGC TGCCCAGC TGCCCAGC TGCCCAGC TCCGACTTC TCCGGAGA GACTGGCC GGAACTGC CGCACTGC GGAGCTGGC CTACACGC GGAGCTGGC CCCCTTTCC GACTGCC CCCCTTTCC GACTGGCC CCCCTGCTCC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGCC CACTGCCG CCCCTGCTCC CACTGCCGC CCCCTGCTCC CACTGCCC CCCTGCTCC CACTGCCC CCCTGCTCC CCCTCTCC CCCTCTC CCCTCTC CCCTCTC CCCTCTCC CCCTCTC CCCTC CCCTCTC CCCTC CCCTCTC CCCTC	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTCC CAGGGTCC CTGGCCGAC ATCAAGAATG TCTTCTTCG GTGTGACA ATCAAGAATG TCTTCTTCTCG GCCAGTGCC CACTGCCGA ACCAGTACC CTGGCCGC CACTGCCGA ATCAGAATA GCTACCGT ATCAGAATA GCTACCGT CCGGCAGC CCTCTGTTG TGGACCC CACTCCGCAA CAGCCCCGCA ACCCAGTACC CAGCCGCC ACCCAGTACC CGGGAGTCACC CGGGAGTCACC CAGCCCCTCCAA CAAGCCACCG CGGAGTCACC CGGGAGTCACC CGGACTCACCT CGGCACCC CGGGAGTCACC CGGGAGTCACC CGGGAGTCACC CGGGAGTCACC CGGAGTCACC CGGGACTCACC CGGAGTCACC CGGGACTCACC CGGAGTCACC CGGGACTCACC CGGAGTCACC CGGAGTCACC CGGAGTCACC CGGAGTCACC CGGAGTCACC CGGACTCACC CGGAGTCACC CGGACC CGGAGTCACC CGGACTCCC CGCACTCCC CGCACC CGCACTCC CGCACC CGCACC CGCACC CGCACC CCCTCCC CGCACC CCCTCC CGCACC CCCTCC CGCACC CCCTCC CGCACC CCCTCC CGCACC CCCTCC CCCCTCC CCCC CC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGTGACCTGA TCAATGACTT CTGTGACTG GCTGACCGA GCACAGGCCA TGACATGAC CTGTGACCGA GCACAGCCA TGACATGAC CTGTGACGG CTGAGGGC CTGAGGCCA TGACATGCC TGACATGCCA AGCAGACCCT TGCTGAGGG CTCAGGGCC AGCAGACCCT TGCTGAGGC AGCAGACCCT TGCTGAGGC CCTGTGAGGC CCTGTGAGGC CCTGTGAGGC CCTGTGAGGC CCTGTGAGGC CCGTGGTTCC AGCTGCCGG CCATCATTGT CAGCATTCGA GAGTGGGTCC	GCAGGGGGGA GCAGCCCAG GCAGCCCAG CTTACTGGAT CGAAGGCTGA CTTACTGAT CGAAGGCTGA TGATTCTC CAAGGTCTGC CAAGGCTGAG GCAGCTGAAG GCAGCTGAG CAGCATCTTG CGTGCCTGTG GCAGCCAAGC CTACAAGGTCAG CAGCTCCTG GGTCCTCTG GGTCCTCAGT CCTGCGTGAG CAGCTCCTC CCTGGCTCACCTC CACTGCCTCAC CACTGGCCACC CCACTGGCCACC CCTGGACCA CCACTGGACCA CCACTGGACCA CCACTGGACCA CCACTGGACCA CCTTGGACGA CCCTTGGACGA CCCTTGGACGA CCCTTGGACGA CCTTGGACGAC CCCTGAGGGG CCCCTGGACGAC CCCTGGACGAC CCGTGAGGGC CCTTGGACGAC CCGTGACGAC CCGTGACGAC CCTTGGACGAC CCGTGACGAC CCGTGACGAC CCTTGGACGAC CCGTGACGAC CCGTGACAC CCGTGACGAC CCGTGACAC CCGTGACGAC CCGTGACAC CCGTGAC CCGTGAC CCGTCAC CCGTCAC CCGTCAC CCGTCAC CCGTCAC CCGTCAC CCCCC CCCCC CCCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ACCACTGCCT AGGACACTAC ACCCGACCTC CAGGACACCTC CAGTACACC GTGCCATCC CAGTACACC GTGCCATCC GTCCAGCTG CTGACCTC GTGACCTTC GTGACCTTG ACCACTGCC GTGGCCTGC GTGGCCTGC GTGGACTTGC CAGTACCTC CAGTACCTC CAGTACCTC GTGACCTTC CAGTACCTC CAGTACCTC GTGGCCTGC GTGGCCTGC GTGGCCTGC CGGCCACTT ACCTCTGCC CGGTTGGACT CAGGTTGCCT CAGGGGGTTC CAGGGGGTTC CAGGGGGTT CAGGCTTG CGAGTGCCTG AGTGCCAGTG AGCACCACTAC ACCACCACCACCACCACCACCACCACCACCACCACCA	120 180 240 300 360 420 480 540 600 660 720 1080 1140 1200 1320 1380 1440 1500 1680 1740 1560 1680 1740 1860
505560657075	Nucleic According sequences of the control of the c	Id Accession lence: 99-89 11 GGCGCTGGGC CTCTAGCTGA CGCGGGATC CACGGGCTT CAGCAATTTC CAGCAATTTC CATCAGCGGC GTTCCAGAG GTCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGAC TGCGGCTT TGCGGGGG GCTGGGCCACAG GCTGGGCCACAG GGAGCTGGC CGCACAGC GGAGCTGGC CGCACAGC CGCACTGC CCCCACTGC CCCCGGGGC CCCGGGGC CCCGGGGC CCCGGGGGC CCCCGGGGGC CCCCCGGGGAG CCCCCGGGGAG	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGTCC CTGGCCGAC ATCAGACACTC CTGGCCGAC ATCAGACACTC CTGGCCGAC CCACTGCCGA CCACTGCCGA ACCACTGCCGA CCACTGCCGA CCACTGCGCAC ATCAGACATA CGCTACCGT CCACTGCCGA ACCCCGCAC CAGGCCCGCC CACCGCCAC CAGGCCCGCC CACCGCCAC CCACTCCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTTTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCTA TCAATGACTT CTGCTGGTG TGATGACTG GTGAGCGGC GTGAGCGGCA GGCACTGAC GTGAGCGGCA TGACTGCC TGAGCGGCA TGACATGGCG TTCAGTGTT TATTTGGCCG TGACAGCCCT TGCCTGAGGC AGAAGGCCT TGCCTGAGGC AGAAGGCCT TGCCTGAGGC CCGTGGTTCC AGCATCATTGT CAGCATTCCT GAGCATTCCT GAGCATTCCT GAGCATTCCT CAGCATTCCT CAGCATTCGTGT CAGCATTCGTGT CACCATTGCTGTCC CACTTGCTGTT CACCATTGCTGTCC CACTTGCTGTT CACCATTCGTGTCC CACTTGCTGTT CACCATTCGTGTCC CACTTGCTGTT CACCATTCGTTCC	GCAGGGGGGA GACCTGCGG GCAGCCCAG CTTACTGGAT CGAAGGCTGC AGGCCAG CTTACTGGAT CGAAGGCTGA AGGCTGAG CAATTCTC CAAGGTCTGC AAGGCTGAAG CAGCATCTTG CGAGCTGAAG CAGCATCTTG CGAGCTGAAG CAGCATCTTG CGAGCTGAAG CACATCTTG CGAGCTGAAG CACATCTTG CGAGCTGAAG CACATCTTG CGAGCTGAAG CACATCTTG CGAGGTGAAC CAGCTCCTG GCTCCTCAGT CCTCCACACT CCTCACACT CCTCACACT CACTGGACCA GCAGCGGTC CCTTGGATGAC CCCTTGGATGAC CCCTTGGAGGC CCCTTGGAGGGC TCCAGGGCTC CCCTGAGGGCT CCCTTGAGGGC CCCTTGAGGGC CCCTTGAGGGC TCCAGGGCTC	GCAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGGG GGAGTTGCCT AGGACACTAC ACCCGACCTC AGCCAATCCT AGCCAATCCT AGCCAATCCT AGCCAATCCT AGCCAATCCT CAGTACACTC GTGCCAGCTG GTGACTGGAC CTGGGCCCACT ATCCTGGGCCCA CTGGGCCATCT CAGTACACTCC CGGTTGGAAT GATGTGAAC GAGTGTCCT CAGGGGTTGCC CGGTTGCCT CAGGGGGTTGCC CGGGTGTCCT CAGGGGGTTG CCGGCTTC CGGGGGTTG CCGGCTTC CGGGGGTTG CCGGGTTGTGG CCGGTTGTGG CCGGTTGTGG CCGGTTGTG CGGGTTGTGG CCGGTTGTGG CCAGTGTCCT CCGGGTTGTGG CCGGTTGTGG CCGGTTGTGG CCAGTGTCT CCGGGTTGTGG CCGGTTGTGG CCAGTGTCCT CCGGGTTGTGG CCGGTTGTGG CCAGTGTCG CCAGTGT CCAGTG CCAGT CCAGTG CCAGTG CCAGTG CCAGTG CCAGTG CCAGTG CCAGTG CCAGTG CCAGT CCAGTG CCAGT CCAGTG CCAGTG CCAGTG CCAGTG CCAGTG CCAGT CCAGTG CCAGTG CCAGT CCAGTG CCAGTG CCAGT CCCAGT CCAGT CC	120 180 240 360 420 480 540 660 720 840 900 1020 1140 1200 1320 1380 1440 1500 1560 1680 1740 1860 1920
505560657075	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGATT CAGCATTTC AGCAATTTC AGCAGACT CTACAAGGG CTTACAAGGG CTGCCCAG GTCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCGGACTT CTGCGCCTG GTGGCCACAGC GGACTTGCT GGGACAGGC GGAACTGAC AGGGCTGGCCACAGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CGCACTGC CCCACTGC CCCCACTC CCCCACTC CCCCACTC CCCCACTC CCCCACTC CCCCACTC CCCCACTC CCCCCCC CCCCCC CCCCCCC CCCCCCC CCCCCC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGGAGGTCTC CTGGCAGAGC GATGCACTTC GTGTGACACTC CTGGCCGAC CTGGTGACACTC GCAGAGACC CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAAGACC CCACTGCCGA ATCAGAGACC CCACTGCCGA ATCAGAGACC CCACTGCCGA ATCAGAGACC CCACTGCCGA ATCAGAGACC CCCTTCTTT GGAACTTC GGGCACGC CACTCGCGCA ACCCTGCAC ACCCCTGCAA ACCCAGTACC CGGAGACCC CCGGAAACTC CCGGAAACTC AGGGTGCCCT AGGGTCAGA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TGACCTTGA TGACCTTG GTGTGTGT CTGTGGTG GGTGCCTGA GGCACAGCCCA GGCACTGAC CTGTAGACTG GTTAGACTG GTTAGACTG GTTAGACTG GTTAGACTG GTTAGACTG GTTAGTGT TATTTGGCCG AGCAGACCCA TGCCTGAGGT CCGTGGTTCC AGCAGACCCG AGCAGCCCG CCATCATTGT CAGTGTTCCG GCATCATTGT CAGCATTCGA GAGTGGTCC CCACTTGCTGT CAGCATCGTGT CAGCATCGTGT CAGCATCGTGT CAGCATCGTGT CAGCATCGTGT CAGCATCGTGT CAGCATCCGT GGGGGACCCGT	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCTGA AGTGCAGTAC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG CGACTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTCAAA GACAGCTCGAG GCAGCTCCAG GCTGCCTCTG GCTGCGTGAC CAGTGTGGCG CAGTGTGCCTCTC CCTCACATC CCTCACACTC CACTGGCCTCT CCTCACACTC CACTGGACCA CCTGGACCA CCTGGAGCAC CCTTGGATGAC CCTTGGAGCAC CCTTGGAGCAC CCTTGGAGCC CCTTGGAGCC CCTTGGAGCC CCCTGAGGCC CCCTGGAGCC CCTGGAGCC CCTTGGAGCC CCTTGGAGCC CCTTGGAGCC CCTTGGAGCC CCTTGGAGCC CCTGGAGCC CCTTGGAGCC CCTTGGAGC CCTTGGAGCC CCTTGGAGCC CCTTGGAGCC CCTTGGACC CCTTGGACC CCTTGGACC CCTTGGACC CCTTGCAC CCTTCC CCTTC	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGAC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGCGGC GGAGTTGCCT AGCGACCTC AGCCATCCC AGCCATCCC GAGTACACTC GTGCCAGCTG GTGCCAGCTG GTGACTGCA GTGACTGCCC GTGGCCTGGC GTGGCCTGGC GTGGCCTGCC CGGTTGGAT ATCCTGGGC CGGTTGGAT GATGTGACCC CAGCTGCCC CGGTTGGAT CCCGCCCACTT ATCCTGGCC CGGTTGGAT GATGTCACCC CAGCTGCCT CAGGGGTTGCAT CAGGGGTTG CAGGGGTTG CAGGGGTTG CAGGGGGTTG CAGGGGGTTG CCGGTTGTGA ACTCAGCCT CAGGGGGTTG CAGGGGGTTG CAGGGGGTTG CCGGTTGTGGA AGTGCCATC CAGGGGGTTG CCGGTTGTGG AGTGCCATC CGGGTTGTGG AGTGCCATTC CGGGTTGTGG AGTGCCATTC CGGGTTGTGG AGTGCCATTC CGGGTTGTGG AGTGCCATTC CGGGTTGTGG AGTGCCATTC CGGGTTGTGG AGTGGATTTC	120 180 240 300 360 420 600 660 720 780 840 900 1020 11200 1260 1320 1320 1340 1500 1680 1740 1860 1740 1860 1920 1980
50556065707580	Nucleic According sequences of the control of the c	id Accession lence: 99-85 11 GCCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCAATTTC AGCCAGTGC CTACAAGGGG CTTACAAGGGG CTGCCCAG GTCCCAGGAC TGCTGTGGGC TGCCCAGGAC TGCCGGACTT CTGCGGACTT CTGCGCAGAC GCCGACTTC GTGGACAGGC GGAACTGAC GGAGCTGGC TGGGGCTGGC TGGGCTGCC AGGTGCCC AACAGCC GAAGGGGGCTG CGCACTGCC CACTGTTCC CACTGGTCC AACAGACCTC CACTGGTCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CCCCGGGGA AACGGAGTGGCC CCCCCGGGGA AACGGAGTGGC CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC CACCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC GAGCACAGGC CGAGCACAGGC CCCGGGGG CCCGGGGG CCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC CGAGCACAGGC CCACGGGGG CCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGCGGGG CCCCGCGGGG CCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCC	#: NM_0000 33 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGAGTCC CAGGAGTCC CAGGACACTC CTGGCAGAC CTGGCAGAC TTCTTCTCTCG GTGTGCACGA ATCAGAGACC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAGACC GCTTCTTCT GCTGGCAGG ATCCAGAATA CCAGATGCC CACTGCCGA ACCAGTGCCC CCTTCTTTG CCTGGCAGC CACTGCCGC CACTGCCAG CCTCGCAC CCTGCAC CCCGCACCC CAGCCGCCC CACCCGCACCC CACCCGCACCC CACCCGCACCC CCGGAACTC CCGGACC CCC CCTC CCGC CCTC CCGC CCC CCC CCC C	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC CTGTGTGCCAC CTGACCTGA CTGACCTGA TCAATGACTT CTGCTGGTGC TGGTGTCCC AGCTGCCAA TCAATGACTT CTGCTGGTGC GTGACCTGA GGTGACCTGAC GTGACCTGAC GTGACATGAC TGACATGAC TGACATGAC TGACATGCC AGCAGACCCA AGCAGACCCT AGCAGACCCT AGCAGACCCT AGCAGACCCT AGCAGACCCT AGCAGACCCT AGCAGACCCT CAGTGTTCC CCGTGGTTCC CCGTGGTTCC CCCTTGCCCG GCATCATTGT CAGCATTCGT GGGGACCCGT AGTCCAGCCA AGTCCGCAC AGTCGCTCT AGCACTCCAC AGTTCCTGT CGGGACCCGT AGTCCACCCA AGCCCAC AGC	GCAGGGGGGA GACGCTGCGG GCAGGCCAG CTTACTGGAT CGAAGGCCTG CGAGGCCTGA GCAATTCTC CAAGGTCTGC AAGGCTGAGG GCAGCTGAGG GCAGCTGAGG CAGCATCTTG CGTGCCTGTG GCAGCCAGGC CAGCATCCTC GCAGTGCCTCTG GCTGCGTGAC CAGCATCCTCT GCTGCGTGAC CAGCTCCTG GCTGCGTGAC CAGTGTGGGG CAGCCCCCTC CCTCACACTC CCTCACACTC CACTGGACCA GCAGCGGCC CTTGGATGAC CCTTGGACCA CCAGGGCT CCCTGGAGGC CCCTTGGATGAC GCCCTGGAGCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC GACACTGCCC CTTGGATGCC GACACTGCCC GACACTGCCC GACACTCCC GACACTCC CTCACACTC CACTGCCC CTTGGATGCC GACACTCC GACACTCC GACACTCC GACACTC CACTGCC CTCGACC CTCTGCC CTCTCACACT CACTC CACTGCC CTCTCACACT CACTC CAC	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA GCGCATCC CATGTGGCTG ATCCTGATCA GCGCACCTC AGCCACCTC AGCCACCTC AGCCAACCTC AGCCAACCTC AGCCAACTC AGCCAACTC AGCCAACTC AGCCAACTC AGCCAACTC AGCCAACTC ACCCTGCCC GTGGCCTGG GTGGCCTGG CTGGCCCACTT ATCCTGGGC CGGTTGGAAT GATGTGACC TACACTCTCC CAGCTGCCTG CAGCTGCCTG CAGCTGCCTG CAGCTGCCTG CAGCTGCTT CAGGGCTTGC CAGCTGCTT CAGGGGTTTC CAGGGGTTTC CAGGGGTTTC CCGGTTTTC CCAGACTCTA	120 180 240 300 360 420 600 660 720 780 840 900 1020 1140 1260 1320 1380 1440 1560 1560 1620 1680 1740 1800 1860 1980 2040
50556065707580	Nucleic According sequences of the control of the c	id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCTT CAGCATTTC CAGCAGTTCT CAGCAGTGCA GTTCGGCCTG CTACAAGGG GTCCCAGGA GTCCCAGGA TGCTGGGC TGCTGGGC TGCTGGGC TGCTGGGC TGCGACTTC TGCGGAGA GACTCTGGC GGAACTGAC GGAGCTGGC GGAACTGAC CGCAGCAC GGAGCTGGC CGCACAGC GGAGCTGGC CGCACAGC CGCGGGGC CGCGGGGC CCCCGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCTGGTGC CCCCGGGGC CCCCGGGGG AACAGACCTG CCCCCGGGGC CCCCCGGGGG CCGCCGGGGG CCGCCG	#: NM_0000 333 21 1 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGCC CAGGGTGCC CTGGCCGAC ATCAAGAATG TCTTCTTCG GTGACACGC CCACTGCGA ATCAAGAATG TCTTCTTCG GCCAGTGCC CACTGCGA ATCAGAGC CCACTGCGA ATCAGAATA GCTACCGG ATCCAGAATA GCTACCGG ATCCAGAATA GCTACCGG CCTCTGTTG TGGACCCC CACTCGCCAA CCACTCCCAA CAAGCCACCC ACCCAGTACC CAGGTGCCC CAGCGCC ACCCAGTACC CGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAACCC CTGCAGCCCC CTGCAGCCCC CCTCCACC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCACCCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCGCAACCCC CCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA TCAATGACTT CTGCTGGTG TCACCTGA TCAATGACTT CTGCTGGTGC GTGAGCGGC GTGAGCGGC GTGAGCGGC GTGAGCGGC AGACATGAC CTGTGAGCGG GTTCAGTGT TATTTGGCC AGCAGACCT AGCAGACCT AGCAGACCC AGCAGACCC AGAGACCCT CCGTGGTTC CAGCTGCCGAG CCACAGCCCA AGCAGACCC CCGTGGTTC CAGCTGCCCG GCATCATTGT CAGCATTCGA GAGTGGGC CACTTGCTGT GGGGACCCGT AGCTCAGCCCA AGCTCGCCG AGCACCCTA AGCTCGCCG AGACCCCTA	GCAGGGGGGA GACGCTGCGG GCAGCCCAG CTTACTGGAT CGAAGGCTGA CGAAGGCTG TGATTCTC CAAGGTCTGC TGCATTCTC CAAGGTCTGA GAGCTGAAG GCAGCTGAAG GCAGCTGAAG CAGCATCTTG CGTGCCTGTG CTACAAGGTC CGAGGTGAAC CAGCTCAGT CCAGCTCCTG GGTCCTCAGT CCTCACACT CCTCACACT CCTCACACT CCTCACACT CCTCAGAGCC CCTGGACCA CCAGGTGCC CCTGGACCA CCAGGTGCC CCTGGACCA CCTGGACCA CCCTGGACCA CCCTGGACCC CCCAGGTGCC CCAGGTGCC CCAGGTGC CCAGGTC CCAGC CCAGCT CCAGGTC CCAGC CCA	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA ACCACTGCCT AGGACACTAC ACCCGACCTC CAGTACACTC GTGCCAGCTG AGCCATCC CAGTACACTC GTGCCAGCTG GTGACTCTC GTGACTCTGA ACCACTGCC GTGGCCTGC GTGGCCTGC GTGGCCTGC GTGGCCTG CCGGCCACTT ATCCTGGGCC CGGTTGGAT CACTGCC GAGTGTGCCTG CGGCTGCC GAGTGTGCCTG CGGCTGCCTG CGGCTTGCC CGGCTTGCACTC CGGGTTGCAT CCGGCCTG CGGCTTGCC CGGTTGCAT CCGGCCTG CCGCCCACTT CAGGCGTTC CCGGCCTG CCGCCACTT CCGGCCTGC CCGCCCACTT CCGGCCTTC CCGGCTTGCCTG CCGGTTGCTG CCGGCTTGCTG CCGGTTGTGT CCGGACTCTA CCGGCTTTT CCGGACTCTTA CTGTCCGTTAC CTGTCCGTTAC	120 180 240 300 360 420 600 660 720 780 840 900 1020 11200 1260 1320 1320 1340 1500 1680 1740 1860 1740 1860 1920 1980
505560657075	Nucleic According sequences of the control of the c	id Accession lence: 99-85 11 GCCGCTGGGC TCCTAGCTGA CGCGGGATC CACGCGCTT CAGCAATTTC AGCCAGTGC CTACAAGGGG CTTACAAGGGG CTGCCCAG GTCCCAGGAC TGCTGTGGGC TGCCCAGGAC TGCCGGACTT CTGCGGACTT CTGCGCAGAC GCCGACTTC GTGGACAGGC GGAACTGAC GGAGCTGGC TGGGGCTGGC TGGGCTGCC AGGTGCCC AACAGCC GAAGGGGGCTG CGCACTGCC CACTGTTCC CACTGGTCC AACAGACCTC CACTGGTCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CACTGGTGCC CCCCGGGGA AACGGAGTGGCC CCCCCGGGGA AACGGAGTGGC CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC CACCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC GAGCACAGGC CGAGCACAGGC CCCGGGGG CCCGGGGG CCCGGGGG GAGCACAGGC GAGCACAGGC GAGCACAGGC CGAGCACAGGC CCACGGGGG CCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGGGGG CCCCGCGGGG CCCCGCGGGG CCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCGCGGGG CCCCCC	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTGCC CAGGGTGCC CTGGCCGAC ATCAGAAATG TCTTCTTCG GCAACACTC CTGGCACAC CTGGTGCAC ATCAGAATG CCACTGCCGA CCACTGCCAA CCACTGCCAA CAGGCCACCC CAGGAACTC CCGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAAACTC CCGGAACCCC CTGCTGCCCCC CCTGCTGCAC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCACCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTG TGATGACCTG TGATGACCTG TGATGACCTG TGATGACCTG TGATGACCTG TGATGACCTG TGACCTGAC TGACATGAC TGACATGCC AGCACCCA TGACATGCC TGACAGCCCA TGACATGCC TGACAGCCCA TGACATGCC TGACAGCCCA TGACATGCC TGACAGCCCA TGACATCGT TATTTGGCC AGCAGCCCA CCATGGTTC CAGCATTCCAGCCA GCATCATTGT CAGCATTCGT GGGGACCCTT AGTCCAGCCA AGACCCCTA TCATCCTGGG CAACCCCTA TCATCCTGGC CAACCCCTA TCATCCTGGC TCATCCTGGC TCATCCTGCCCTA TCATCCTGGC TCATCCTGCCCTA TCATCCTGGC TCACCCCCTA TCATCCTGGC TCACCCCCTA TCATCCTGCCCTACCCCCTA TCATCCTGCCCTACCCCCTA TCATCCTGCCCTACCCCCTA TCATCCTGCCCCTACCCCCCTACCCCCCTACCCCCCTACCCCCCTACCCCCC	GCAGGGGGGA GACGCTGCGG GCAGCCCAG CTTACTGGAT CGAAGGCTGC AGGCCAG CTTACTGGAT CGAAGGCTGC AGGCTGAG GCAGCTGAG GCAGCTGAG CAGCATCTTG CGAGCTGAG CAGCATCTTG CGAGCTGAG CAGCATCTTG CGAGCTGAG CAGCATCTTG CGAGCTGAG CAGCATCTTG GGAGCTGAAG CCACGTGCCTGTG GCAGCTCCAG CAGCCTCCAG CAGCCTCCTG GCTCCTCAG CCAGTGGGC CCCTGGACCC CCTGGACCC CTTGGATGC CCCTGGAGCC CCCTGGAGCC CCCAGGGCTC CCCAGGGCTC CCCAGGGCTC CCCTGGAGCC CCCAGGGCTC CCCAGGGCTC CCCAGGGCTC CCCAGGGCTC CCCAGGGCTC CCCAGGGCCC CCCAGGGCCC CCCAGGGCCC CCCAGGGCCC CCCAGGGCCC CCCAGGTGCC CCCAGGTGCC CCCAGGTGCC CCCAGGTGCC TCCAACCGCC TTCGAACGGC TCCAACCGCC TTCGAACGGCC TCCAACCGCC TTCGAACGGCC TTCGAACGGCC TTCGAACGGCC TTCGAACGGCC	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATTCC CATGTGGCTG ATCCTGATCA ACCACTGCCT AGGACACTAC ACCCGACCTC AGCCAATCCT AGGACACTCC GTGCCATCC GTGCCATCC GTGCCATCC GTGCCACTC GTGGCCTG GTGGCCTG CCGCACTT ATCCTGGGCC CGGTTGGAC GTGGCCTG CCGCACTT CAGTACCTC CGGCACTT CAGTACCTC CGGCACTT CAGTGCCC CGGTTGGAC CCGCACTT CAGGACTCC CGGTTGCC CGGTTGCC CGGTTGCC CGGTTGCC CGGTTGCC CGGTTGTCC CGGGTTCCC CGGTTGTGC CCGGCTTC CGGGGTTC CCGGCTCT CAGGGGTTC CCGGTTGTGG AGTGCCAGTG CCGGTTGTGG AGTGCCAGTG CCGGTTGTGG AGTGCATTC CCAGCTCTA CCCACTGGCC CCACTGGCC CCACTGCC CCACTGGCC CCACTGGCC CCACTGGCC CCACTGGCC CCACTGCC CCACTGC CCACTGCC CCACTGCC CCACTGCC CCACTGCC CCACTGC CCACTGCC CCACTGC CCACTC CCACT	120 180 240 300 360 420 540 660 720 840 900 900 1020 1140 1260 1380 1440 1500 1560 1620 1680 1740 1860 1860 1920
50556065707580	Nucleic According sequences of the control of the c	Id Accession lence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGGCTT CAGCAATTTC CAGCAATTTC CATCAAGGG CTTCCAGAC GTCCCAGGA TTCCGGAGA TTCCGGAGA CTCCGACTTC TTCCCGGAGA GACCTCTGCT TTCCCGGAGA GACCTCTGCT GGACTGCC AGGACTGC CCCACACAC GGAGCTGGCC AGGACTGC CCCACACAC CCCCACACAC CCCCACACAC CCCCCACAC CCCCCGGGCC CCCCGGGCC CCCCGGGCC CCCCGGGGC CACAGACCTC CCCCGGGGCC CCCCGGGGA AAGGCACAGC CCCCCGGGAG AAGGCACAGC CACCACACAGC CCCCCGGGAG AAGGCACAGC CACCACACAGC CACCACACAGC CCCCCGGGAG AAGGCACAGGC CACACACA	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGAGTGCC CAGGGTGTGC CTGGCCGAC ATCAGAATG TCTTCTTCTC GCAGAGGC CCACTGCCGA ACCAGAGAC CCACTGCCGA ACCAGAGAC CGCAGTTCC CTGGCCGAC ATCAGGAGTC CCACTGCCGA ACCAGAGAC CCACTGCCGA ACCAGAGAC CCACTGCCGA CCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCGA CCCACTGCCA ACCCCTGCAA CCCAGTACC CGGAGACTC CCGGAAACTC AGGGTGCCT AGTGGTCCGC CCGCAGACCCC CCCGCAAACCCC CCCGCAAACCCC CCCGCAAACCCC CCCGCAAACCCC CCCGCAAACCCC CCCGCAAACCCC CCCGCAACCCC CCCGCAACCCC CCCGCAACCCC CCCGCAACCCC CCCGCAACCCC CCCCTCCAC ACCCCTGCCAC CCCCTCCACCCC CCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGCCAC GCTTTGCCAC CTGACCCTGA TCAATGACTT CTGCTGGTG TCAATGACTT CTGTGGTG TGGTGCCA GGCACAGCCGA GGCACAGCCGA GGCACAGCCGA GGCACAGCCGA TGACTTGCTGACTGC GTTAATTGCCG GTTAATTGCCG GTTCAGTGT TATTTGGCCG AGCAGACCCT TGCCTGAGGG CCACTGAC CCGTGGTTCC AGCAGCCCA AGCTGCCCG GCATCATTGT CAGCATTCGT CAGTTCCAGCCA AGCTGCCCG GCATCATTGT CAGCATTCGT CACTTGCTGT CAGCACCC GAACCACCTA TCATCGTGGC GCACCCAT CCACTTGCTGCCCG GCACCCCC GCACCCCCC GCACCCCC GCACCCCCC GCACCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCCC GCACCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCC GCACCCCCCC GCACCCCCC GCACCCCCC GCACCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCC GCACCCCCCCC	GCAGGGGGGA GACGCTGCGG GCAGGCCCAG CTTACTGGAT CGAAGGCCTAC CGAAGGCTGAC CGAAGTCTC CAAGGTCTGC AAGGCTGAG GCAGCTGAG CAGCATCTTG CGTGCCTGTG GCAGCTCAA GCACCTCAA GCACCTCAG GCAGCTCCAG GCACCTCCTG GCTCCTCACAT CCATGGCCTACT CCTCACACTC CACTGGCCTCT CCTCACACTC CACTGGACC CCTTGGATGAC CCGTGAGGCC CCTTGGATGAC CCGTGAGGCC CCTTGGATGAC CCCTTGGATGAC CCCTGGAGCC CCAGGTGCCT CCCCAGGGCT CCCCAGGGCT CCCCTGGAGCC CACTGGACCA CCCAGGTGGCT CCCAGGGCT CCCAGGGCC CCAGGTGGCT CCCAGGGCC CAGGTGGCT TCCAACGGC TCCAACGGC CCAGGTGGCT TCCAACGGC TCCACACT TCCACCAT TCCACCA	GCAAGGGACA CTTCTGGTGG GCACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATCAC CGCGCCATTCC CATGTGGCTG ATCCTGATCA ACCCGACCTC AGCCAATCAC ACCCGACCTC AGCCAATCAC ACCCGACCTC GTGCCATCT AGCACTGCC GTGGCCATCC CAGTTGGAC ACCAGCTC CAGTCGGCCACTT ATCCTGGGCC GTGGCCTGGACT GATGTGGACT CCGGCCACTT CCGGCCACTT CCGGCCACTT CCGGCCACT CCGGCCACTT CCGGCCACTT CCGGCCACTT CCGGCCACTT CCGGCTGGACT CCGGTTGGAAT GATGTGACC CGGTTGGATT CCAGGGGTTG CCGGTTGTGCAC CCGGTTGTGA ACCTCTCC CGGTTGCACT CCGGTTGCCAC CCGGTTGGACT CCGGTTGTGA CCGGTTGTGCCAC CCGGTTGTGG AGTGCCAT CCGGTTGTGG AGTGCCAT CCGGTTGTGG AGTGCCAT CCACTGGGCC ACCTGGGCC ACCTGGACCA	120 180 240 360 420 480 540 660 720 840 900 1020 1140 1260 1320 1380 1440 1560 1680 1740 1860 1920 1980 2010 2160

		700 077 3	GCCACGGTGG	omes conces	TO CA CTOCA C	CCACATACTC	2340
			GCCACGGTGG				2400
	AGTATACGGT	GCATGTGAGG	CCCCATGIGG	CIGGCGIGGA	CCTCCACATC	CTCAATGCTT	2460
	TTGTGAGGAC	TGCCCCTGAG	CCTGTGGGTC	GIGIGICGAG	ACCCACACATC	TACAGACTCC	2520
5			ACCTGGGTAG				-
J	CCTGGGGCCG	GAGTGAAGGC	GCCCCATGA	GGCACCAGAT	ACTCCCAGGA	AACACAGACT	2580
			GAAGGTGGAG				2640
	TCGGGGACCG	CGAGGGCACA	CCTGTCTCCA	TTGTTGTCAC	TACGCCGCCT	GAGGCTCCGC	2700
	CAGCCCTGGG	GACGCTTCAC	GTGGTGCAGC	GCGGGGAGCA	CTCGCTGAGG	CTGCGCTGGG	2760
10			GGCTTCCTTC				2820
10	AGTCCCGGGT	CCTGGGGCCC	GAGCTCAGCA	GCTATCACCT	GGACGGGCTG	GAGCCAGCGA	2880
	CACAGTACCG	CGTGAGGCTG	AGTGTCCTAG	GCCCGCTGG	AGAAGGCCC	TCTGCAGAGG	2940
			CCTCGTGTTC				3000
			GCCTGGACTC				3060
			CCTGGCCAGG				3120
15	GGATCTCAAG	CTCCCAGCGG	GTGACAGGGC	TAGAGCCTGG	CGTCTCTTAC	ATCTTCTCCC	3180
	TGACGCCTGT	CCTGGATGGT	GTGCGGGGTC	CTGAGGCATC	TGTCACACAG	ACGCCAGTGT	3240
	GCCCCCGTGG	CCTGGCGGAT	GTGGTGTTCC	TACCACATGC	CACTCAAGAC	AATGCTCACC	3300
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	JOHNSHOOT		.21.1001101				
		79 Protein cession #: (
80	1	11	21	31	41	51	
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00							
80		82 DNA sequ				-	
		id Accession Lence: 44-54	1 #: BC00129	1			
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                                                                               2100
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		97 Protein cession #: 1			•		
10	KRIVAHAVEV	PAVQSPRRSP	21 SPTKPLRRSQ RISFFLEKEN	EPPGRELTKE	DLFKTHSVPA	TPTSTPVPNP	60 120
15		VPRVCAKPWA	KVRRSYSRLE PDMTLPGISP				180 240
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	1	11	21	31	41	51 	
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	CCAATAATTG	CAAAACAGCT	CTTTAGCAGC	TTGTTTTCTG	GAATTTTGAA	AGAGATGGAT	8520
	מים מים מים מים מים מים	CVCACACACA	סגמסגממממ	ልጥሮልሮጥሮልልል	AGTITGCTTCA	AGACTTCAAT	8580
80	WOULT TANGE	ADDICTOR		TOTAL COMMENT	TOTAL TOTAL	TCACCACAM	0640
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	ACRECCOCCC		COMOUNT WAGE	TO A TOTOGO	AATACOACO	CCTCCGTGGG	8880
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236

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       RSPTRGSEFS GKGDVPNTSL NSTSQPVTKL ATEKDISLTS QTVTELPPHT VEGTSASLND
55
       GSKTVLRSPH MNLSGTAESL NTVSITEYEE ESLLTSPKLD TGAEDSSGSS PATSAIPFIS
       ENISQGYIFS SENPETITYD VLIPESARNA SEDSTSSGSE ESLKDPSMEG NVWFPSSTDI
       TAQPDVGSGR ESPLQTNYTE IRVDESEKTT KSFSAGPVMS QGPSVTDLEM PHYSTFAYFP
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       TEVTPHAFTP SSRQQDLVST VNVVYSQTTQ PVYNGETPLQ PSYSSEVFPL VTPLLLDNQI
                                                                           780
       LNTTPAASSS DSALHATPVF PSVDVSFESI LSSYDGAPLL PFSSASFSSE LFRHLHTVSQ
                                                                           840
60
                                                                           900
       ILPOVISATE SDKVPLHASL PVAGGDLLLE PSLAQYSDVL STTHAASETL EFGSESGVLY
       KTLMFSOVEP PSSDAMMHAR SSGPEPSYAL SDNEGSQHIF TVSYSSAIPV HDSVGVTYQG
                                                                           960
       SLPSGPSHIP IPKSSLITPT ASLLQPTHAL SGDGEWSGAS SDSEPLLPDT DGLTALNISS
                                                                          1020
       PVSVAEFTYT TSVFGDDNKA LSKSEIIYGN ETELQIPSFN EMVYPSESTV MPNMYDNVNK
                                                                          1080
       LNASLQETSV SISSTKGMFP GSLAHTTTKV FDHEISQVPE NNFSVQPTHT VSQASGDTSL
                                                                          1140
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       KPVLSANSEP ASSDPASSEM LSPSTQLLFY ETSASFSTEV LLQPSFQASD VDTLLKTVLP
                                                                          1200
       AVPSDPILVE TPKVDKISST MLHLIVSNSA SSENMLHSTS VPVFDVSPTS HMHSASLQGL
                                                                          1260
       TISYASEKYE PVLLKSESSH QVVPSLYSND ELFQTANLEI NQAHPPKGRH VFATPVLSID
                                                                          1320
       EPLNTLINKL IHSDEILTST KSSVTGKVFA GIPTVASDTF VSTDHSVPIG NGHVAITAVS
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       PHRDGSVTST KLLFPSKATS ELSHSAKSDA GLVGGGEDGD TDDDGDDDDD DRGSDGLSIH
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70
       KCMSCSSYRE SQEKVMNDSD THENSLMDQN NPISYSLSEN SEEDNRVTSV SSDSQTGMDR
                                                                          1500
       SPGKSPSANG LSQKHNDGKE ENDIQTGSAL LPLSPESKAW AVLTSDEESG SGQGTSDSLN
                                                                          1560
       ENETSTOPSF ADTNEKDADG ILAAGDSEIT PGFPQSPTSS VTSENSEVFH VSEAEASNSS
                                                                          1620
       HESRIGLAEG LESEKKAVIP LVIVSALTPI CLVVLVGILI YWRKCFQTAH FYLEDSTSPR
                                                                          1680
       VISTPPTPIP PISDDVGAIP IKHFPKHVAD LHASSGFTEE FETLKEFYQE VQSCTVDLGI
                                                                          1740
75
       TADSSNHPDN KHKNRYINIV AYDHSRVKLA QLAEKDGKLT DYINANYVDG YNRPKAYIAA
                                                                          1800
       QGPLKSTAED FWRMIWEHNV EVIVMITNLV EKGRRKCDQY WPADGSEEYG NPLVTQKSVQ
                                                                          1860
       VLAYYTVRNF TLRNTKIKKG SQKGRPSGRV VTQYHYTQWP DMGVPEYSLP VLTFVRKAAY
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       AKRHAVGPVV VHCSAGVGRT GTYIVLDSML QQIQHEGTVN IFGFLKHIRS QRNYLVQTEE
       QYVPIHDTLV EAILSKETEV LDSHIHAYVN ALLIPGPAGK TKLEKQPQLL SQSNIQQSDY
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       SAALKQCNRE KNRTSSIIPV ERSRVGISSL SGEGTDYINA SYIMGYYQSN EFIITQHPLL
80
                                                                          2100
       HTIKDFWRMI WDHNAOLVVM IPDGONMAED EFVYWPNKDE PINCESFKVT LMAEEHKCLS
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       VHDEHGGVTA GTPCALTTLM HQLEKENSVD VYQVAKMINL MRPGVFADIE QYQFLYKVIL
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       SLVSTROEEN PSTSLDSNGA ALPDGNIAES LESLV
85
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Seq ID NO: 181 DNA sequence Nucleic Acid Accession #: Eos sequence

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3			CTCACTTCGA				60 120
			TCTGGAAATG				180
			CCTGGATTGG				240
10			GTCCTATACA				300
10			CCCAAAACAA GAAACTTAAA				360 420
			TGGGAAAACA				480
			AATGGTGTTT				540
1.5			TGGATCAGAG				600
15			TGATGCGGAC				660
			TTTATCCATT TGGAGTCGAA				720 780
			GAACCTTCTG				B40
~^	AATGGCTCAT	TGACATCTCC	TCCCTGCACA	GACACAGTTG	ACTGGATTGT	TTTTAAAGAT	900
20			CCAGTTGGCT				960
			GGACTACTTA CTCATACACT				1020 1080
			TCAGGCTGAC				1140
0.5			TTATGATACC				1200
25			AACCAAGCAT				1260
			GCTACCCAAT AAAATACAGC				1320 1380
			CCCTGAATTA				1440
••			AGAAGGCGCT				1500
30			ACCCCAGATT				1560
			GACTAACCGA ATCTTTAAAT				1620
			GACTTCTCAG				1680 1740
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			GCTTGATACT				1920
			CATCTCTGAG ATATGATGTC				1980 2040
			TTCAGAAGAA				2100
40	GTGTGGTTTC	CTAGCTCTAC	AGACATAACA	GCACAGCCCG	ATGTTGGATC	AGGCAGAGAG	2160
			CACTGAGATA				2220
			GATGTCACAG CTTCCCAACT				2280 2340
			CTCCACGGTC				2400
45	GTATACAATG	CAGAGGCCAG	TAATAGTAGC	CATGAGTCTC	GTATTGGTCT	AGCTGAGGGG	2460
			AGTTATACCC				2520
			TATTCTCATC ATCCCCTAGA				2580 2640
			AGCAATTCCA				2700
50	TTACATGCAA	GTAGTGGGTT	TACTGAAGAA	TTTGAGACAC	TGAAAGAGTT	TTACCAGGAA	2760
			CTTAGGTATT				2820
			AAATATCGTT CAAACTGACT				2880 2940
			TATTGCTGCC				3000
55			ACATAATGTG				3060
			TGATCAGTAC				3120
			GAGTGTGCAA AAAAAAGGGC				3180 3240
			GCAGTGGCCT				3300
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						CAGTATGTTG	
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						CTATGTTAAT	
65						CCAGCTCCTG	
						CAACAGGGAA	
*						TTCATCCCTG CCAGAGCAAT	3780
							3900
70							3960
						TAAGGTCACT	
						TCAGGACTTT GTGTCCTAAA	
						TATAAAAGAA	
75						AGTGACGGCA	
							4320
						TGACATTGAG GGAAGAGAAT	4380 4440
00						AGCTGAGAGC	
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						TTTCCCATCA	
						TAACAATGTG TTGAATTTTA	4680
						CAGAAAATTT	
85	CAATTTATAG	AGGTTAGGAA	TTCCAAACTA	CAGAAAATGT	TTGTTTTTAG	TGTCAAATTT	4860
•						TACAGTAGCC	
	TGTAAATAAA	ACACTCTTCC	ATATGATATT	CAACATTTTA	CAACIGCAGT	ATTCACCTAA	4980

5	AGTAGAAATA ATATTTATAA ATTGTTTAGT GTATTGTGTT ATAGAAATAC TCAAATGGTT	TTGTAGATTT TTAATGACGT ACCTAAGTCA CTTCATTTTG	TATTGTAAAT TTATATTTTA AGTTCATTAG TTAACTTTGT AAAGAAGTTT GAATTGCAAA AAAAAAA	CTACTGAGTC CTGGTCTTAC TTCAGCATGT TTATGAGAAT	AAGTITTCTA TCTACCAGTT AATTTTAACT AACACCTTAC	GTTCTGTGTA TTCTGACATT TTTGTGGAAA CAAACATTGT	5040 5100 5160 5220 5280 5340
10		182 Protein cession #: 1	a sequence: Eos sequence	•			
	1	11	21	31	41	51	
15	MDIT KDET VO	I OT I CACCATA	 WANGYYRQQR	 	TCALNOVNWG	 	60
13			KFQGWDKTSL				120
	PKASKITFHW	GKCNMSSDGS	EHSLEGOKFP	LEMQIYCFDA	DRFSSFEEAV	KGKGKLRALS	180
			ESVSRFGKQA				240
20			AVFCEVLTMQ DPENYTSLLV				300 360
20			NMSYVLQIVA				420
			AIVNPGRDSA				480
			NSTSQPVTKL				540 600
25			NTVSITEYEE VLIPESARNA				660
	TAOPDVGSGR	ESPLOTNYTE	IRVDESEKTT	KSFSAGPVMS	QGPSVTDLEM	PHYSTFAYFP	720
	TEVTPHAFTP	SSRQQDLVST	VNVVYSQTTQ	PVYNAEASNS	SHESRIGLAE	GLESEKKAVI	780
			IYWRKCFQTA EFETLKEFYQ				840 900
30			TDYINANYVD				960
	VEVIVMITNL	VEKGRRKCDQ	YWPADGSEEY	GNFLVTQKSV	QVLAYYTVRN	FTLRNTKIKK	1020
			PDMGVPEYSL				1080 1140
			NIFGFLKHIR KTKLEKQFQL				1200
35	VERSRVGISS	LSGEGTDYIN	ASYIMGYYQS	NEFIITOHPL	LHTIKDFWRM	IWDHNAQLVV	1260
			EPINCESFKV				1320
			KTFELISVIK LMRPGVFADI				1380 1440
40	AALPDGNIAE						
40	Nucleic Aci	183 DNA sec d Accession mence: 148-4	#: EOS sec	quence			
45	1	11	21	31	41	51	
45	1	11 	21 	31 	1	1	
45	CACACATACG	 CACGCACGAT	CTCACTTCGA	 TCTATACACT	 GGAGGATTAA	AACAAACAAA	60 120
	CACACATACG CAAAAAAAAAC	CACGCACGAT ATTTCCTTCG	CTCACTTCGA CTCCCCCTCC	 TCTATACACT CTCTCCACTC	 GGAGGATTAA TGAGAAGCAG	AACAAACAAA AGGAGCCGCA	60 120 180
45 50	CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG	CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCTGGATTGG	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA	120 180 240
	CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG	CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG	120 180 240 300
	CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG	CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCTGGATTGG	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA	120 180 240
50	CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTGT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC	CTCACTTCGA CTCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAACCAA GAACTTAAA TGGGAAAACA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA	AACAAACAAA AGGAGCCCCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT	120 180 240 300 360 420 480
	CACACATACG CAAAAAAAA CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCATAACAC GAGTTTCAGA	CTCACTTCGA CTCCCCCTCC TCTGGAATTG CCTGGATTGG GTCCTATACA CCCAAAACAA GAAACTTAAA TGGGAAAACA AATGGTGTTT	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT	AACAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA	120 180 240 300 360 420 480 540
50	CACACATACG CAAAAAAAAC CAGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATC AACACATTCA ACACACTGAG AAATGCAATA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAG GAGTTTCAGA TGTCATCTGA	CTCACTTCGA CTCCCCTCC CTCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACAA GAAACTTAAA TGGGAAAACA AATGGTGTTT TGGATCAGAG	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GCAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGGACAAAA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT	120 180 240 300 360 420 480
50	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAAA GGAAAAGGGA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTCATCTGA ACTCACTGCTT AGTTAAGAGC	CTCACTTCGA CTCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA TGGGAAACA AATGGTGTTT TGGATCAGA TGATCCAGT TTTTCCATT	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTACA TGTTTCAAG TTGTTTCAAG TTGTTTGAGG	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAA GTTTTGAGGA TTGGGACAGA	ACCAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGGAAAATTTG	120 180 240 300 360 420 480 540 600 660 720
50 55	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAAA GGAAAAGGGA GAATTCAAAG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTCATCTGA TCTACTGCTT AGTTAAGAGC CGATTATTGA	CTCACTTCGA CTCCCCTCC TCTGGAATG CCTGGATTGG GTCCTATACA CCCAAAACAA TGGGAAAACA AATGGTGTTT TGGATCAGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTCAA TGTTTCAA TGTTTCAA ATGTTTCAA AGTGTTTGAG AGTGTTAGTC	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGAGGA GTTTTGGGAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGCAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAAA AGAAAATTTG GCAGGCTGCT	120 180 240 300 360 420 480 540 600 660 720 780
50	CACACATACG CAAAAAAAA CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAATA GAGATGCAATA GAGAAAGGGA GAAAAGGGA TTAGATCCAT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TTCATAACAC GAGTTTCAGA TGTCATCTGA TCTACTGCTT AGTTAAGAC CGATTATAGA TCATACTGT TCATACTGT	CTCACTTCGA CTCCCCTCC TCTGGAATG CCTGGATTGG GTCCTATACA CCCAAAACAA TGGGAAAACA AATGGTGTTT TGGATCAGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTTA AAAGCAAGCA CATAGTTTAG CGATTTTCAG TTGTTTGAGG AGTGTTAGTC CCAAACTCAA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAA ATATTGATGA ATCACTAA AGATAACTTA AGGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGAGGA GTTTTGGGACAGA GTTTTGGGACA CTGACAAGTA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC	120 180 240 300 360 420 480 540 600 660 720 780
50 55	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCCAATA GGAAAAGGGA GATTTCAAAG GTTTCAAAG TTAGATCCAT AATGGCTCAT AACACTTAGCA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATCAGA TCTACTGGTT AGTTAAGAGC CGATTATGA TCAATCGTT TGACATCTGT TGACATCTCC TCTCTGAAAG	CTCACTTCGA CTCACTTCGA CTCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AGGACTTAAA AGGGAAACA AATGGTGTTT TGGATCAGAC TTTATCCATT TGGAGTCGAA GAACCTTCTG GAACCTTCTG CCCTGCACA CCAGTTGGCT	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTT AAGGACAAA GGTTTTGAGGA TTGGGACAGA GTTTTGGGAA GTTTTGGGAA ACTGGACAGA ACTGGACAGT ACTGGACAGT ACTGGACTGA	ACCAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGGAATTTGCACTT TTACATTTG TTACATTTAC TTTTTAAAGAT AATGCAACAA	120 180 240 300 360 420 480 540 660 720 780 840 900 960
50 55	CACACATACG CAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GTCAGCGGAG AAATGCAATCA GAAATGCAAA GGAAAAGGGA GATTTCAAAG TTAGATCCAT AATGCTCAT AATGCTCAT ACAGTTAGAC TTAGATCCAT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTCATCTGA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATACTGCT TGACATCTCC TCTCTGAAAG TCATGCTGAT	CTCACTTCGA CTCCCCTCC CTCGCATTGG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGATCAGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA GAACCTTCTG TCCCTGCAC CCCTGCAC GGACTACTTA	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA GTGGAAATTA AAAGCAAGCA CGATTTTCAA CGATTTTCAA TTGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACAGTTA GACACAGTTA GTTTTTTTTTT	GGAGGATTAA TGAGAAGCA AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGAGAA GTTTTGGGAAA CTGACTAA ACTGATTAT ACTGGATTAT ACTGGATTAT ACTGGATTAT ACTGGATTAT ACTGGATTAT ACTGGATTAT ACTGGATGAT TTCGAGAGCA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAAAGAT TTTTAAAGAT AATGCAACAA ACAGTACAAG	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020
50 55 60	CACACATACG CAAAAAAAA CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAATA GAGATGCAATA GAGATGCAATA AATGCATCAT ATTGATCCAT AATGGCTCAT ACAGTTAGAT TCTGGTTATG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAAATAG TCATAAACAC GAGTTTCAGA TGTCATCTGA TCTACTGCTT TAGTTAAGAGC CGATTATTAGA TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGT TCATCTGTT TGACATCTCC TCTCTGAAAA AGGTGTTTTC	CTCACTTCGA CTCCCCTCC TCTGGAATG CCTGGATTGG GTCCTATACA CCCAAAACAA GAAACTTAAA TGGGAAAACA AATGGTGTTT TGGATCAGAG TGATCCAGAT TGAGTCCAT TGGAGTCGAA GAACCTTCTG CCCTGCACA CCAGTTGGCT CGACTACTTA CGCTACTT CGCTACACT CGCTACACT CCCTACACA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG CGATTTTCAGG AGTGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA AGGATAAACTTT AAGGACAAAAC ATTTGAGAGA GTTTTGAGGA GTTTTGAGGA GTTTTGAGGA CTGACAAGTA ACTGGATTGT AAGTTCTTG AAGTTCTTGA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTAAAGAT AATGCAACAA ACAGTACAAG ACAGTACAAG AGCAGTTTGT	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020
50 55	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GATTTCAAAG GTTAGATCCAT AATGGCTCAT AATGGCTCAT ACAGTTAGAT TCTGGTTATG TCTCTTAGAC AGTTCAGAAC	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCAATAACAC GAGTTCACAGA TCTACTGCTA AGTTAATAGAC CGATTATGA TCAACTGT TCAACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCACACTCTC TCCTCGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTCAGGTCGT	CTCACTTCGA CTCACTTCGA CTCACTCC TCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AGAGCTTAAA TGGGAAAACA AATGGTGTTT TGGATCAGAC TTTATCCATT TGGAGTCGAA GAACCTTCTA CCCTGCACA CCAGTTGGCT GCACTACTTA CTCATACCTT TCAGGCTGAC TCATGCACA TTATGCATAC CTCATACACT TCAGGCTGAC TTATGATACC TTATGATACC	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCA AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTTT AAGGACAAA GTTTTGAGGA TTGGGACAGA GTTTTGGGAA GTTTTGGGAA GTTTTGGGTAA ACTGGATTGT AAGTTCTTAC TTCGAGAGCA AGATTCTTAC TTCGAGAGCA AGATTCATGA	ACCAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT TACAGTCAAA AGGAGTCAAA AGGAGTCAAT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTTTCTTTTTACA TCTTGTTACA TTTGTACCAG	120 180 240 300 480 540 600 660 720 780 960 1020 1020 1140 1200
50 55 60	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAAA GGAAAAGGGA CATTCAAAG TTAGATCCAT ACAGTTAGAC TCTGGTTATG TCTCTTAGAC TCTGGTTATG TCTCTTAGAC CAGTTCAGAAC CGGAAAAGC CAGTTGGATG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCATAACAC GAGTTCATGA TCTATAACAC GAGTTATAGA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTGAGAGACCCA GAGAGAACCCA	CTCACTTCGA CTCCCCTCC CTCGCATTGG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGGATAGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA GAACTTCTG TCCTGCACA CCAGTTGCT TCAGGCTAC TCAGGCTTAC TCAGGCTTAC TATGATACACT TCAGGCTTAC TATGATACC AACCAAGCAT	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GACACGTTG GTTTTTTGTG CAAAACAATT GGAAAGGAAG CCAGAGAATT ATGATTGAG AGATTGAG AGATTGAG AGATTGTGAG AGATTTTTTGTG CAAAACAATT GGAAAGGAAG CCAGAGAATT ATGATTGAG AAATTTTTGA	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACC ATCTACACTAA AGATAACTTT AAGGACAAAC TTGGGACAGA GTTTTGGGAA CTGACAAGTA ACTGGATTGT AAGTTCTTAC TTCGAGAGCA AGATTCATGA AGATTCATGA AGATTCATGA AGATTCATGA AGATTCATGA ATACCAGCCT CAGATGGCTA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGA ATTTCCACTT TACAGTAAA AGAAAATTTG GCAGGCTGCT TTACATTTAA TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCGTTTGTTACA TTGTTACA TTTTGTTACAG TCTTGTTACCAG TCTAGAGCTTG	120 180 240 360 420 480 540 660 720 780 840 960 1020 1080 1140 1200 1260
50556065	CACACATACG CAAAAAAAA CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAATA GAGATGCAATA AATGCAATA AATGCTCAT AATGGCTCAT ACAGTTAGAC TTCTCTAGAC TTCTAGAC TTCTCTAGAC AGTTCAGAC TTGGATCAGAC TTGGATAGAC GGTTCAGAC GGTTCAGAC GGTTCAGAC GGTTCAGAC GGTTCAGAC GGTTCAGTT GGGTAAGAC GGTTCATTC	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG CAGATTCAGA TTCATAACAC GAGTTTCAGA TGTAATAGAGC CGATTAATAGAC CGATTAATAGAC CGATTATTGA TCATACTGCTT TGACATCTCC TCTCGAAAA AGGTGTTTTC CAGAAAATGT CTGAGTCGT GAGAGGACCA GAGAGGACCA TCAATAATTT	CTCACTTCGA CTCCCCTCC CTCGCATTGG CTCGGATTGG GTCCTATACA CCCAAAACAA AAACATTAAA TGGGAAACAA AATGGTGTTT TGGATCAGAG TTATCCATT TGGATCCATT TGGAGTCGAA CCAGTTGGCT GCACTACTAC CTCATACACT TCAGGCTGAC TCATACACT TCAGGCTGAC TATGATACACT TCAGGCTGAC TATGATACACT TCAGGCTGAC TATGATACACT TCAGGCTGAC TATGATACACT TCAGCCAACCA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG GAATTTTAGG AGTTTTGAGG AGTGTTAGTC CCAAACTCAA GACACGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA AGGATAACTTA AGGATAACTTT AAGGACAAAAC ATTTGAGGA GTTTTGAGGA GTTTTGAGGA CTGACAAGTA ACTGGATTGT AAGTTCTTG AAGTTCTTG AAGTTCTTGA AGATTCTTGAGAA ATACCAGCCT AGTTTTGAGGA ATACCAGCCT AGTTTTGAGGT ATTTTGAGT ATTTTGAGT ATTTTTAGGATT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTTGT TCTTGTTACA TTTGTACCAG TCTTGTTACA TTTGTACCAG AGCAGTTGT TCTTGTTACA ATGTACAAG AGCAGTTTGT TCTTGTTACA ATGTACCAG AGCAGTTTGT TCAAGACTTG AGTAGCCATA	120 180 240 300 480 540 600 660 720 780 960 1020 1020 1140 1200
50 55 60	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAATA GGAAAAGGGA GATTTCAAAG GTTAGATCCAT AATGGCTATA AATGGCTATA ACTGGTTATG TCTGGTTATG TCTCTAGAC TGGGAAAGGCA CGGTTGGATC CGGTTGTATC TGGGAAAGAC CAGTTCGAAC CAGTTCGATATC TGCACTATC TGCACTATC TGCACTAATG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCATAACAC GAGTTCACAGA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGC TCATCTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCATCACTGCT TCAGAAAATGT CAGAAAATGT CTCAGAGTCGT GAGAGGACCA TCAATAATTTT GCTTATATGG	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGATCAGA TGATGCAGAC TTTATCCATT TGGAGTCGAA GAACCTTCTG TCCCTGCACA CCAGTTGGCT TCATGCATT TCAGGCTGAC TTATGATACC TCATACACT TCAGGCTGAC TTATGATACC AACCAAGCAT GCTACCCAAT CCAATACCC AACCAAGCAT CCTACCCAAT AAAATACAGC CCCTGAATTA	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAGACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCA AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTT AAGGACAAA GGTTTTGAGGA TTGGGACAGA GTTTTGAGGA ACTGTACAAAAAA ACTGTTTAGAGAA ACTGTTTTAGAGAA ACTGTTTTACACAACTT AAGTTCTTAC TTCGAGAGCA AGATTCATGA AGATTCATGA AGATTCATGA TTCTTCAGGT AGTTTTGCACT TTCTTCACAT TTCTTCACAT TTCTTCACAT TTCTTCACAT AAGAAATAAT	ACCAACCAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGAA ATTTCCACTT AGCAGTCAAA AGGAGTCAAA AGAAATTTG CAGGCTGCT TTACAATTTAC ATTTTAAAGAT AATGCAACAA ACAGTACCAAG AGCAGTTCTGT TCTTGTTACA TTTTGTACCAG TCAAGACTTG AGTAGCCATTG AGTAGCCATTG AGTAGCCATTG AGTAGCCATTG AGTAGCCATTG AGTAGCCATTG AGTAGCATTG AGTAGCATG AGCATAGACT CAAGGAGGAG	120 180 240 300 360 420 480 540 660 720 780 900 960 1020 1140 1260 11320 1440
50556065	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GATTTCAAAG TTAGATCCAT ACAGTTAGATC TCTGGTTATG TCTCTTAGAC ACTTCATAG TCTCTTAGAC ACTTCAGAC ACTTCAGAC ACTTCAGAC ACTTCAGAC AGTTCAGAC AGTTCAGAT CAGTTCAGAC AGTTCAGAAC AGTTCAGAAC AGTTCAGAAC AGTTCAGAAC AGTTCAGAAC AGTTCACAAC AGTTCAGAAC AGTTCAGAAC AATCCTGAAC GAAGAGGAA	CACGCACGAT ATTTCCTTCG CAGGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATGAT AGTCATCTGA AGTTAATGG TGATATTGA TCATACTGCTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTGAGAGCCCA TCAGTGGT CAGAGACCA TCAATATTTT GCTTATATTG CTTATATTG TCAGATCTTTT AGGACATTTTT AAGACATTTTT AAGACATTTTA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGGATACA AATGGTGTTT TGGATCAGA GTACCATT TGGATCAGA CCAGTTCTG TCCTGCACA CCAGTTGCT TCAGTACACT TCAGTTGCT TCATACACT TCAGTACACT TCAGTACACA CACCACAC AACAACAT CTCATACACT ACCACAC CACTCACAC CACTCACT	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GACACGATT GGAAACAATT GGAAAGAAT ATGATTGTG CCAAACTTAG GCCAGGAATT GGAAAGGAAG ATGATTGAG ATGATTTTGA ATGATTTGAG ATGAGTATTT	GGAGGATTAA TGAGAAGCA ATCAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGGATAACTT AAGGACAAC ATCTCACTAA AGATAACTT AAGGACAAGTA ATTTGAGGA TTTGAGGA GTTTTGGGAA ACTGGATTGT AAGTTCTTAC TCGAGAGCA AGATTCATGA ATACCAGCT AGATTGCAGAT TTGTCAGAT TTGTCAGAT TTGTCAGAT TTGTCAGAT TTGTCAGAT TTGTCAGAT TTGTCAGAT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGGA ATTTCCACTT TCACTGGGAA AGAAATTTG GCAGGCTGCT TTACATTTAA AGAAATTTG ATTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTTGT TCTTGTTACA TCTTGTTACA TCTTGTTACA GCCTACTGAT GCCTACTGAT CCAGGAGGAG CCAGGAGGAG CAGGGCTACA	120 180 240 360 420 480 540 660 720 780 840 960 1020 1140 1200 1140 1320 1320 1340 1500
50556065	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GTCAGCGGAG AAATGCAATA GAAATGCAAA GGAAAAGGGA GATTTCAAAG TTAGATCCAT AATGCTCAT ACAGTTAGAC TTAGATCCAT ACAGTTAGCA TCTGGTATG TCTCTAGAC AGTTCAGAAC TGGAAAAGGGA GAGTGCAAT GGGAAAAGC GAGTGATG AATCCTGAAC GAGAGGGAA AACCAAATCA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTCATCTGA TCATACTGCTT AGATACTGTT TGACATCTCT TCATACTGCTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTCGAGTCGT TCAGAGTCGT TCAGAGTCGT TCAGATCGT AGACATTGA GGAAAAAGGA	CTCACTTCGA CTCCCCTCC CTCGCATTGG CTCGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGATCAGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA GAACCTTCTG TCCTGCACA CCAGTTGGCT CCCTGCACA CAGTTGCT CTCATACACT TCAGGCTGAC TATGATAC CTATGATAC CTATGATAC CTATGATAC CTATGATAC CTATGATAC CTATGATAC CTATGATAC CTATGATAC CCTGACTA AACAAGCAT AAAATACAGC CCCTGAATT AGAAGGCGCT ACCCCAGATT ACCCCAGATT	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA GTGGAAATTA AAAGCAAGCA CATAGTTTCAA GCATTTCAA GCATTTCAA GCATTTCAA GCAACTCAA GACACAGTTG GTTTTTTGTG CAAAACAATT GGAAAGGAAG ATGATTTGTG ATGATTTTTGTG ATGATTTTTTGTG ATGATTTTTTGTG ATGATTTTTTGTG ATGATTTTTTGTG ATTGATTTTTTTGTG ATTGATTTTTTGTG ATTGATTTTTTTGTG ATTGAACTGA ATTGGAACTGA ATTGGAACTG ATTGGAACT CTCACCACAA	GGAGGATTAA TGAGAACAG ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCACATAA AGATAACTTT AAGGACAAAAC GTTTTGAGA GTTTTGAGA GTTTTGAGA CTGACAAGTA ACTGATTAC ACTGATTAC ACTGATTAC ACTGATTAC ACTGATTAC ACTGATTAC ATACCAGCCT AGATGACTA ACTTCAGAT TTCTCAGAT TCTGTAGAGA CACACTACAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTTTAC TTTTAAAGAT TATTGCAACAA ACAGTACAAG AGCAGTTTGT TCTGTTTAC TCTTGTTTACA TCTTGTTTACA TCTTGTTACAG TCAAGACTTG AGTAGCCATA GCCTACTGAT CAAGGAGCAG CAGTGCTACA TCAAGGAGGAG CAGTGCTACA TCGCATAGGG	120 180 240 360 420 480 540 660 720 900 900 900 1080 1140 1260 1380 1440 1500 1500
5055606570	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GATTTCAAAG GTTAGATCCAT ACAGTTAGATCAT ACAGTTAGAC TCTGGTATG TCTCTTAGAC TCTGGTATG TCTCTTAGAC TGGAAAAGAC CAGTTGGATG TTCCTAGAAC TGGAAAAGAC CAGTTGGATG TTCCTGAAC CAGTTCAGAAC AGGTGCTATTC TGCACTAATG AATCCTGAAC GAAGAGGGAA AACCAAATCA ACGAAATCA AAGGAGAGTGT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATCTGCTT TGACATCTCC TCTCTGAAAG TCATGCTGAT TCAGCTGAT TCAGAGAAATGT CCAGAAAATGT CCTGAAAAATGT TCATATTGG GGAAAAATGT TCATATTGG TCAGATCGTT AGGAGACCA TCAATAATTT AGGACATCTT AAGACATTGA GGAAAAATGA TTGATCTTTT AAGACATTGA TTGATCTTTT AAGACCAA TTCCCAATAACTTA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTCC CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGATCAGAC TTATCCATT TGGAGTCGAA GAACCTTCTG TCCCTGCACA CCAGTTGGCT TCAGGCTGAC TTATGATACC TCATACACT TCAGGCTGAC CTATACACT TCAGGCTGC CCAGATTA CCCAAACAT CCTAACACAT CCTAACCAAT CCTAACCAAT CCTAACCAAT CCTGAATTA AGAAGGCGCT ACCCCAGATT AGAAGGCGCT ACCCCAGATT AGCAGACCGA ATCTTTAAAT	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GCAAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GCACAGTTG GCAAACACATT AGAACAATT AGAACAATT AGAATTTGAA ATGATTTGA ATGATTTGAA ATGATTTGA ATGATTTGA ATGATTTGA ATGATTTGA ATGATTTGA ATGATTTGA ATTGACACAC ATTGGAACT ATTGTACACACA TCCCCAACACA TCCCCAACAC TCCCCAACACA	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTTT AAGGACAACTTTGAGGACAGT TTGGGACAGT ACTGGACAGT ACTGGACAGT ACTGGACAGT ACTGGACAGT ACTGCACAGT ACTGCACAGT ACTGCACAGT ACTGCACAGT ACTTTCACGACACT AGATTCATCA AGATTCACACT AGATGGCAT AAGAAATAAT CTGGTAGAGA AACCACTACAA AACCACTACAA AAGCAGTCAC	ACCAACCAAA AGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA AGACTATCGGAA ATCATTGGAA ATCATTGGAA ATCATTGCACT TCACTGT TCACTGT TTACAATTTG AGAGATCTAA AGAAATTTG GCAGGCTGCT TTACAATTTA ACAGTACAAG ACAGTACAAA ACAGTACAAG AGCAGTTCTA TCTTGTTACA TCTTGTACCAG TCAAGACTTG AGTAGCCATA ACAGTACCAAT CCAAGACTTG AGTAGCCATA CCAAGACTTG AGTACCAAG TCAAGACTTG AGTAGCATAC ACCCTACTGAAT CCAAGACTTG CAAGGAGGAG CCTACTGAAT CCAAGACTTGAAT CCAAGACTTACTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CAAGACTTCTCTGAAT CCAAGACTTCTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTGAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTCTAAT CCAAGACTTTCTCTCTAAT CCAAGACTTTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTCTAAT CCAAGACTTCTCTCTCTCTCTCTCTCTCTTCTTCTTCTTCTTCTT	120 180 240 300 360 420 480 660 720 780 960 1020 1140 1200 1140 1200 1140 1500 1500 1620 1680
50556065	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA ACACATTCA GTCAGCGGAG GATGCAAA GAGATGCAAT GAGATGCAAT AATGCATTA ACAGTTCAAG GATTCAAG TTAGATCCAT ACAGTTAGAC TTTCTTAGAC TTCTCTAGAC TCTGGTTATG TCTCTTAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATACA	CACGCACGAT ATTTCCTTCG CAGGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATGAT AGTTAATAGAC GAGTTAATGA TCTACTGCTT AGTAATGAT TGACATCTCA TCATACTGTT TGACATCTCC TCATACTGTT TGACATCTCC TCATGCAAAA TCTACTGTT GCTAATACTTT GCTAATACTTT GCTTATATGG TCATATATGG TCAGATCATTT AGAGCCAT AGAACATTT AAGACCAT ATGAAGCCA ATGAAGCACA ATGAAGCCAA ATGAAGCCAA ATGAAGCCAA ATGAAGCCAA ATGAAGCCAATAC	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTCC TCTGGAAATG CCTGGATTGG GTCCTATACA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCGAGA CTTATCCATT TGGAGTCGAA GAACCTTCTG GAACCTTCTG GACTACTAC CCAGTTGGCT TCAGGCTGAC TTATGCATA CTCATACCAT TCAGGCTGAC CCAGTTACCAAT AAATACAC CCCTGAATTA AGAAGCGCT ACCCAGATTT ACCCAGATTT ACCATAACCT ACCAGACTT ACCACAGACT ACCACAGACT ACCACAGACT ACCACAGACT ACCACAGACT ACCACAGATT ACCCACAGATT ACCCCAGATT ACCCCACACT ACCCACACT ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCCACACT ACCCACACT ACCCACACT ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCTACCCAC ACCTACCCAC ACTTCTCAC CCTCGAATTA ACACCCAC ACCTACCCAC ACCTACCAC ACCTACCAC ACCTACCAC ACCTACCAC ACCTACCAC ACCTACAC ACCTACCAC ACCTACAC ACCTACCAC ACCTACAC ACCTACCAC ACCTACAC ACCTACAC ACCTACCAC ACCTACAC ACCTAC	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAG GGATTTTCAG GGATTTTCAG GCATTTTCAG GCAAACTCAA GACACGTTG GCTTTTTTGTG CAAACAATT GGAAAGAAT ATGATTTGAG ACAGGAAT ATGATTTGAG ACAGGAAT ATGATTAGA ATGATTAGA CCAGAGAATT ATGATTAGA ATGAGTTAG CACCACCAC ATTGTGAACC ACTGCCCAACA TCCACCACAA TCCACCTCCC ACTGTGACTG	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTT AAGGACAAACTTT AAGGACAAACTTT AAGGACAAGTA ACTTTGAGGA GTTTTGAGGA GTTTTGGGAA ACTGGATTGTT AAGTTCTTAC TTCGACAAGTA ACTGGATTCTTAC TTCGAGACC TTCGAGACC TTCTTCAGAT TCGTAGAGA CACACTACAA GAGGAAGTGA AACCAGTCAC AACTGCCAC AACTGCCAC	ACCAACATA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAT TGACTACCGT TCACTGGGA ATTTCCACTT TACACTTACA AGAAAATTTG GCAGGCTGCT TTACATTTAA AGAAAATTTG TTACATTTAA ACAGTACAAA ACAGTACAAA ACAGTACAAG AGCAGTTTGT TCTTGTTACA TTTGTTACA TCTTGTTACA TCTACTGAT CAAGACCTTG AGTAGCCATA CCAGCATACGGA ATCTCTCTGGA ATTCTCTCTGA ATTCTCTCTGGA ATTCTCTCTGGA ATTCTCTCTGGA ATTCTCTCTGGA ATTCTCTCTGGA ATTCTCTCTGGA ATTATTAGCC TCACACTGTG	120 180 240 300 360 420 480 660 720 780 960 1020 1140 1260 1140 1500 1560 1560 1680 1740
5055606570	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GACATGCAATA GGAAAAGGGA TTAGATCCAT ACAGTTAGAC TTTGATCAT ACAGTTAGCA TTTCATAGC TTCTCAGAC TTCTAGAC TTCTAGAC TTCTAGAC TTCTAGAC TTCTAGAC TTCTAGAC TTCTAGAC AGTTCAGAAC CAGTTGGATG GGAAAAGC AATCCTGAAC AACCAAATA AACCAAATAC AAGGGTGATG AACGAAAAAAC AAGGGTGATG AACGAAAAAAC AAGGGTGATG AACGAAAAAC AAGGGTGATG AACGAAAAAC AAGGGTGATG AACGAAAAAC AAGGGTGATG AACTGTCGGG GAAGAAAAAC AAGGGTGATG AACTGTCGGG GAAGAAAAAC AAGGGTGATG AACTGTCGGG	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TCTACTGTT AGTTAAGAGC CGATTATGA TCTACTGTT TGACATCTGT TGACATCTCC TCTCTGAAAG TCATCTGTT AGGTGTTTC CAGAAAATGT CTGAGTGTTT CTGAGTCGT GAGAGACCA TCACATACTGTT CAGAAAATGT CTGAGTCGT GAGAAAATGT CTGAGTCGT TCACATACTTT AGAAAATGT CTCAATACTTT CAGAAAAATGT CTCAATACTTT CAGAAAAATGT CTCAATACTTT CAGAAAAATTT CAGACTCTTT CAGCCAATAC ATATTTCCTTT CAGCCTCTTT CAGCTCTTT CAGCCTCTTT CAGCTCTTT CAGCCTCTTT CAGCTCTTT CAGCCTCTTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCTT CAGCCTCT CAGCCT CAGCC CAGCCT CAGCCT CAGCCT CAGCCT CAGCCT CAGCC CA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCGAG TGATGCAGAC TTTATCCATT TGGAGTCGAA GAACTTCTG TCCCTGCACA CCAGTTGGCT GGACTACTTA CTCATACACT TCAGGCTGAC TATGATACC AACCAAGCAT GCTACCCAATTA AGAAGGCGCT GACTAACCGA ATCTTTAAAT GACTTCTCAG ACCACGAT ACCCCAGATT ACCTCTCAG ATCTTTAAAT GACTTCTCAG AAATGATGGC ATCCTTAAAT	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GCAGACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GCACACGTTG GGAAACAGAT ATGATTTGAG GAAACAGAT GGAAAGAAT ATGATTTGAG AAATTTTGAG AAATTTTGAG AAATTTTGAG AAATTTTGAG AATTTTGAG AATTGAGATTAG CAAACTCAA ATTGGAACTC ATTGGAACTC ATTGGAACTC ACTGTGACTC ACTGTGACTC ACAGTTTCTA	GGAGGATTAA TGAGAAGCA AGCATTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA ATGTTGAGGA TTGGGACAGA GTTTTGGGAA CTGACAAGTA ACTGACAAGTA ACTGACACT AGATTCATCA AGATTCATCA AGATTCATCA AGATTCATCA AGATTCATCA AGATTCATCA AGATTCATCA ATTCTTCAGAT TTGTCGACAT ATGTTCAGAT TTGTCGACAT AGAAATAAT CTGGTAGGAT ACACTACAA CACTACCAA CACTACCAA CACTGCCACC TTCTTAGATT TAACAGATTA	ACCAACAGA AGGAGGAAAA AGGAGCACA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGAA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTGTT TCTTGTTACA TTGTACCAG TCAAGACTTG AGTAGCATA GCCTACTGTAC AGGAGGAGAG CAGTGCTACA TCACAGAGCTTCC CACACTGTG TCACACTTCTGA TCACACTTCTGA TCACACTTCTGA TCACACTTCTGA TCACACTCTGT TCACACTCTGT TCACACTCTGT TCACACTCTGT TCACACTCTGT TCACACTCTGT TCACACTATGT TCACACTCTGT TCACACTATGT TCACACATATGT TCACACATATGT TCACACATATGT TCACACACTCTGT TCACACACTCTT TCACACACTCTGT TCACACACACTCT TCACACACTCT TCACACACTCT TCACACACTCT TCACACACTCT TCACACACTCT TCACACACTCT TCACACACTC TCACACTC TCACACACTC TCACACACTC TCACACACTC TCACACACTC TCACACACTC TCACACACTC TCACACACA	120 180 240 360 420 480 660 720 960 1080 1140 1200 1320 1380 1440 1560 1680 1740 1860
5055606570	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAATA GAGATGCAATA AATGCATTA AATGCATTA AATGCATTA AATGCATTA AATGCATTA AATGCATTA AATGCATTA ACAGTTAGAC TTCTCTAGAC TCTCGTTATG TCTCTTAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC ACGAAATCA AACGAAATCA AACGAAATCA AACGAAATCA AACGAAATCA AACGAAATCA AACGAAATCA AAGGGTGTT AACTTGTCGG ACAGAAAAAG GAAGGTACTT AACTTGTCGG ACTTTGTCGG ACGAAAAAAG GAAGGTACTT AACTTGTCGG ACTTTATTGA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCATAACAC GAGTTCAGAA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATACTGCT TCATCACTGT TGACATCTCC TCATACTGCT TCATCTGCT TCATCTGCT TCATCTGCT TCATCTGCT TCATCTGCT TCATCTGCT TCATCTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCAGACACTCT CAGAAAATGT TCTTATATGG GGAAAAATGT TTGATCTTTT AAGACATTGA GGAAAAAGGA ATGAAGCCAA ATTTTCCTT CGGCTCTTT CAGCCTCTTT CAGCCTCTTA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTCC CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGATCAGA TGATGCAGAC TTTATCCATT TGGAGTCGAA CACATCTCA GAACCTTCTC GGACTACTA CTCATACCAT TCAGGCTGCT TCAGGCTGCC TTATGCATA CTCATACCAT TCAGGCTGC CCTGAATTA AGAAGACT CCCTGAATTA AGAAGACT TACCAAT AGAATACAGC CCCTGAATTA AGAAGACCT ACCCAGATT GACTAACCGA ATCTTTAAAT GACTTCCAG AAATGATGCC AACTACCAA CCCTGAATTA CCTTAAAAT GACTTCTCAG AAATGATGCC AACTTCTCAG AAATGATGCC CTTGATACT CTCTTAAAT CCTTCAACAT CCTCAACAT CCTCAACA	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTCAA TTGTTTGAG AGTGTTAGAG AGTGTTAGTC CCAAACTCAA GCACACTTG GCAAACACAA TTGTTTTGAG GCAGAATTTTTGAG GCAGAACTTG GCAGAACTTAG ATGATTTTGA GAACACAGTTG GACCACTGA ATTGTGAACT ATTGTGAACTG ATTGTGAACT CTCTACCACAA TCCCCAACACA TCCCAACACA TCCACTTCCC ACTGTGACTG CTTAAAACTG CCTAAAACTG ACAGTTCTA GCAGCTGACG ACAGTTCTAA	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACTT AAGGACAAC ATCTCACTAA AGATAACTTT AAGGACAAC ATTTGAGGA TTGGGACAGA GTTTTGAGGA ACTGTTTTACAGAT AAGATTCTTAC TTCGAGAGCA AGATTCTTAC TTCGAGAGCA AGATTCTTAC TTCTCAGAT TTGTTGAGAT TTGTCGACAT AAGAAATAAT CTGGTAGAG CACACTACAA GAGGAAGTGA AACCAGTCAC AACTGCCAC TTCTTAGATT AACCAACTA AACAGACTAC AACTGCCAC TTCTTAGATT CTCAGATT CTCAGATT CTGTAGATC CACTTCAGATT CTCAGATC CACTTCAGATT CTCAGATC CACTTCAGATC CACTTCAGATC CACTTCAGATC CTCATAGATC AACTACAAA	ACCACTOTE TCACACTCC TCACACTCC TCACACTCC TCACACTCC TTATACA ATTATCACAT TCACTGGGA ATTTCCACTT TCACTGGGA ATTTCCACTT TTACATTTC TTACATTTAC ACAGTCAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA TCTTTTACACTTT TCTTGTTACA TTTTTACACTTTAC TCTTGTACA TCTTGTACCAG TCAAGACTTG TCAAGACTTG TCAAGACTACAA ACAGTACAAA ACAGTACCAT TCTTCTTCTGA TCAAGACTACG TCACACTTGT TCACACTTGT TCACACTTGT TCACACTTGT TCACACTTGT TCACACTCCC TCACACTCCC	120 180 240 360 420 480 540 660 720 780 960 1020 11200 11200 11200 1140 1500 1560 1740 1860 1740 1860 1920
505560657075	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAAT AACACATTCA GTCAGCGGAG GAATGCAATA GGAAAAGGGA GATTCAAAG GTTAGATCAT AATGCATTA ACAGTTAGAC TTTGATCAT ACAGTTAGAC TCTGGTTATG TCTCTTAGAC CAGTTCAGAC CAGTTCAGAC CAGTTCAGAC ACGAAATCA AACGGAAATCA ACGAAATCA ACGAAATCTGTCGG GAAGGTACTT AACTTGTCGG GAGGTACTT ACCTTGTCGG GAGGTACTT ACCTTGTCGG GAGGTACTT ACCTTGTCGG GAGGTACTT ACCTTGTCGG GAGGTACTTT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATGAT ACTCATCATA ACTCATACTGCT AGATACTGTT TGACATCTCA TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGCT TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT TCAATACTTT CCTTATATGG TCAATACTTT AGACCTTTT AAGACCATAC ATATTTCCTT CAGCCTCTTT CGACCTCATT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTT CAGCTT CAGCTT CAGCTT CAGCT	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTCC CTCGGAAATG CCTGGATTGG GTCCTATACA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCAGA CTTATCCATT TGGAGTCGAA GAACCTTCTG GACTACTAC CCAGTTGGC TCCTGCACA CCAGTTGGCT TCAGGCTGAC TTATGATACC AACCAGCTT AAAATACAC CCCTGAATTA AGAAGCGCT ACCCAGATT CACTTACAC AATGATGCC ATCCTTAAAT CATTCTCAG AATGATCC CATCCTTAAAT CCTTGATACT CATCTCTGAG	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAAGCACTGA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAA GTTTTCAG GTTTTCAG GTTTTTGAG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTGTG CAAACAATT GGAAAGAAT ATGATTTTGAG ACAGTTAGT CCACACACT ATTGTGAACT ATTGTGAACT ATTGTGAACT ATTGTGAACT CCCCAACAA TCCACCACAA TCCACCTACCC ACTGTGACT CTTAAAACTG ACAGTTTCTA ACAGTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTCTA ACAGTTTCTA ACAGTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTCTA ACAGTT	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAACC ATCTACTAA AGGATAACTT AAGGACAAA GGTTTTCGAGAA GTTTTGAGGA TTGGGACAGA GTTTTGAGGA ATTGATGA ACTGGATTGT AAGTTCTTAC TTCGAGAGCA AGATTCTTAC TTCGAGAGCA AGATTCTTAC TTCTTCAGAT TTGTCGACAT TTGTCGACAT ACTGTAGAGA CAGATGCAT ACAGATGCAT ACAGATGCAT AAGAAATAAT CTGGTAGAGA AACCACAC AACTGCAC AACTGCAC TTCTTAGATC AACTGCAC TTCTTAGATC AACTGCAC AACTGCCAC TTCTTAGATC TAACAGAATTA ATTCTTCAGAT AATTCTTCAGAT AACAGAATAA ATTCTTCAGAG AACTGCCAC TTCTTAGATC AACTGCACC AACTGCCAC AACTCCAC AACTGCCAC AACTCCAC AACTC	ACCACATTC CACACTCC CACACTC CACACT CACACT CACACTC CACACTC CACACTC CACACTC CACACT CACACTC CACACT CACACT CACACT CACACT CA	120 180 240 360 420 480 540 660 720 900 900 1020 1140 1260 1320 1440 1560 1560 1680 1680 1680 1780 1800 1800 1900
5055606570	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA GTCAGCGGAG GAATGCAATA GAGATGCAATA GAGATGCAATA AATGCATATA AATGCATAT AATGCTCAT ACAGTTAGAC TTCTTGGTTATG TTCTTAGAC TCTGGTTATG TTCTTAGAC CAGTTGGATG GGTGCTATTC GGTGCTATTC GATCCTAATG AATCCTGAAC AAGGGGAAAAC AACGAAATCA AAGGGTGAT AACGAAATCA AAGGGTACTT AACTTGTCGG GAAAACCCAG GCAACTTCTG GAAACTCTCTG GAAAACCCAG	CACGCACGAT ATTTCCTTCG CAGGACCCG GTGTTTGCCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATCAGA TCTACTGCTT AGTTAATAGAC CGATTATTGA TCATACTGCTT AGTTAATAGAC CGATTATTGA TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGAGAATTT CAGAAAATTT GCTTATATGG TCAGAGACCA TCAATACTTT GCATACTGT TGACATCTC TCAGAGACCA TCAATACTTT CTGAAAG TCAGACATTT CAGAAAATTT CCTATTT CAGACATCTT CAGCCTCTTT GGACTGCAGA CCAGTTTCAA CTATCCCATT AGACAATAAC CTATCCCATT AGACAATAAC	CTCACTTCGA CTCACTTCGA CTCCCCTCC CTCGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGGATACA AATGGTGTTT TGGATCAGAG GTATCCATT TGGATCAGAG CTTATCCATT TGGAGTCACA CCAGTTCTG GCACTACTTA CTCATACACT TCAGGCTGAC TTATCATACACT TCAGGCTAC CCCTGAATTA AGAAGCAT GCTACCCAAT AAAATACAGC CCCTGAATTA AGAAGGCGCT ACCCAGATT GACTACCCAAT AGAATGATGC CTTATACACT CTCTGAAC ATCTTTAAAT GCTTGATACC AACTAACACT CCCTGAATTA AGAAGGCGCT ACCCCAGATT GACTACCCAAT CCTTGATACAC AATGATGCC ATCTTCACAG AATGATGCC ATCTTGATAC CATCTCTCACG ATTATGATGTC CATCTCTCACG ATTATGATGTC CATCTCTCACG ATTATGATGTC	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTAA AAGCAAGCA CGATTTTCAA TGTTTGAG AGTTTTGAG AGTGTTAGTC CCAAACTCAA GACACAGTT GGAAAGCAATT GGAAAGGAAG ATGTTTTGTG CCAAACTGAA ATGATTTTGAG ATGATTTTGAG ATGATTTTTGA ATGATTGAACT ATGATTGAACTG ATTGGAACTG ATTGGAACTG ATTGGAACTG ATTGGAACTG ATTGGAACT ATCCCCACAA TCCACACAA TCCACACAT TCCACACAC ACTGGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACTGACTG ACTGGACTG ACTGGACTG ACTGGACTG ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCTA ACAGTTTCCC CTTATACCAC	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGATAAACA ATCACATAA GGGATAAAC ATCTCACTAA AGATAACTT AAGGACAAA GTTTTGAGAA GTTTTGAGAA GTTTTGAGAA GTTTTGGGAA CTGACAAGTA ACTGATTAT ACTGATTAT ACTGATTAC TTCGAGAGCA AGATTCATAC AGATTCATCA AGATTCATCA TTCTCAGAT TTCTCAGAT TTCTCAGAT ACTGCACAC CTGATACAA CAGATGCAA AACTGCCAC TTCTTAGATC TAACAGCTA AACTGCACC TTCTTAGATC TAACAGATTA ATTCTTCAGA ATTCTCAGA ATTCTCAGA ATTCTTCAGAT AACTGCACC TTCTTAGATC AACTGCACC ACTGCACC AACTGCACC ACTGCACAC AACTGCACAC AACTGCACAC AACTGCACAC AACTGCACAC AACTGCACC AACTGCACAC AACTCCACAC AACTGCACAC AACTGCACAC AACTGCACAC AACTGCACAC AACTGCACAC AACTACACAC AACTGCACAC AACTGCACAC AACTACACAC AACTACACAC AACTACACAC AACTGCACAC AACTACACAC AACTACAC AACTACACAC AACTACACAC AACTACACAC AACTACACAC AACTACACAC AACTAC	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTGCGT TCACTGGGA ATTTCCACTT TCACTGGGA ATTTCCACTT TTACATTTGA AGGAGATAAA AGGAAAATTTG GCAGGCTGCT TTACATTTAA AGAAATTTG TTTTAAAGAT AATGCACAAA ACAGTACAAG AGCAGTTGT TCTTGTTACA TCTTGTTACA GCCTACTGAT CAAGACCTTG AGTAGCCATA GCCATAGGG TCAAGACTTG TCACAGTACAA TCACACTTGGA TCACACTTGGA TCACACTTGGA TCACACTTGGA TCACACTTGGA TCACACTTGGA TCACACTTGT TCCACACTTGT TCCACACTCTC TCACACTCTCC AAATGCTTCC	120 180 240 360 420 480 540 660 720 840 900 900 1080 1140 1260 1380 1450 1560 1580 1680 1860 1980 1980 2040
505560657075	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GATTTCAAAG GTTAGATCAT ACAGTTAGAT ACAGTTAGCA TCTGGTTATG TCTCTAGAC TCTGGAAAAC AGGTAATG AATCCTGAAC CAGTTCAGAAC AGGTAATG AATCCTGAAC GAAGAGGGAA AACCAAATCA AAGGGTAATG AACCAAATCA AAGGTACTT AACTTGTCGG AGTTTGTCGAC CGAGAAAAAG GAAGGTACTT AACTTGTCGG AGTTTATTGA GCAACTTCTG GAAAACCCAA GGAGATTCTG GAAAACCCAA GGAGATTCTG GAAAACCCAA GCAACTTCTG GAAAACCCAA GTGTGCTCTTC	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTAATGA TCTACTGCTT AGTTAAGAGC CGATTATTGA TCATCTGCTT AGTTAAGAGC CGATTATTGA TCATCTCCT TCTCTGAAAG TCATCTCCT TCTCTGAAAG TCATGCTGAT CAGAAAATGT CTGAGTCGT GAGAGACCA TCAATAATTT AAGACATTTA AGACATTTC TTGATCTTT AAGACATTGA CTATCTCTT CAGCCTCTTT GGACTCGAG CCAGTTCAA CTATCCCATT AGACAATAAC CTTCATCAG CTTACCCATT AGACAATAAC CTTCATCAG CTTACCCATT CAGCCATTCAA CTATCCCATT CAGCCATTCAA CTATCCCATT CAGCCACTAC CTTCATCAGG CTTACTCAC	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTTCG CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AATGGTGTTT TGGGACACA TTATCCATT TGGAGTCGAA CAACCTTCTG CCAGTTGGT TCAGTGCACA CTATACATA CTCATACACT TTATGATAC CTCATACCAT TCAGCCAAT AAAATACAC CCCTGAATTA AGAAGGCGT ACCCCAGATTA AGAAGGCGCT ACCCCAGATTA AGAAGGCGCT ACCCCAGATTA AGAAGGCGCT ACCCCAGATTA AGAAGGCGCT CACTTAAACT CATCTCTCAG AAATAACT CATCTCTCAG AAATGATGCT CATCTCTGAG ATATGATACT CTTCAGAAAAA AGACATAACA	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG GGATTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GCACAGTTG GTTTTTTGTG CAAAACAATT ATGATTGAG AATGATTAGA AATGATTAGA AATGATTAGA AATGATTATGA ATTGAAACTA ATTGAAACTGA ATTGGAACTC ACTGACACCA ACCCACCACACA ACCCACACACA ACCACACTCC CCTAAAACTC ACTGTAAAACTG ACAGTTTCTC CACACTCC ACTGTGACTC CCTAAAACTG ACAGTTTCTC CCACTCC CCTTAAAACTG ACAGTTCCC CTTAAACCAC ACAGTTCCC CTTAAACCAC ACAGTTCCC CTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTTAAACCAC CCTCACACACCCCC CCACCACCCCC CCACCACCCCC CCACCA	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAACTA ATCTCACTAA AGGATAACTT AAGGACAAA ATATTGATGA GGATAACTT AAGGACAAA TTGGGACAGA TTGGGACAGA GTTTTGGGAC AGTTTTGAGGA ATCTCATA AGATCATTA AGGTTCTTAC TTCGAGAGCA AGATTCATGA ATACCAGCCT AGTTTGCAGT TTGTTCAGT AGGTTGTACAT ATGTTGAGAA ACTGCACA ACTCTCAA AACTACCAA AACTACCAAC AACTACCAA AACTACCAAC AACTACCAA AACTACCAAC AACTA	ACCAACTAG AGGAGCACA AGGAGCACA AGGAGCAAA AGGAGCAAA ACGATTCTACA ATCATTGGAA TGACTGGAA ATTTCCACTT ACCAGGGAAAA ATTTCCACTT AGCAGTCAAA AGAAATTTG GCAGGCTGCT TTACATTTAA AGAAATTTG ACTACATAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACCAA TCTTGTTACA TTTGTACCAG TCAAGACTTG AGCAGTACTAA ACAGTACTAA ACAGTACCAAT ACCACTATGA ACCATATG ACCATATG ACCACTAGG CAAGACTTG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTAGG TCACACTATG TCACACTATACACTATACACTATACACTATACACTATACACTATACACTATACACTATACACTATACACTATACACTATACACTACACTACACTACACTACACTACACTACACTACACTACACTACT	120 180 240 300 360 420 480 540 660 720 780 960 1080 1140 1200 1320 1380 1440 1560 1680 1740 1860 1920 1980 2010 2160
505560657075	CACACATACG CAAAAAAAAC CAGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAAT AACACATTCA GTCAGCGGAG AAATGCAATA GAGATGCAAT AATGCATCA TTAGATCCAT ACAGTTAGAC TTAGATCCAT ACAGTTAGAC TCTCGTTATG TCTCTTAGAC TCTCGTTATG TCTCTTAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC GAAAATCA AACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCT ACTTGTCGG AGTTTATTG GAAACCCAG GAAGATTCAG GAAGATTCAG AGTTTGTCG GAAACCCAG GAAGATTCAC GAAGATTCAC GAAGATTCAC GAAGATTCAC GAAACCCAG GAAGATTCAC GAAACCCAG GAAGATTCAC GAAACCCAG GAAGATTCAC GAAACCCAG GAAGATTCAC GAAACCCAG GAAGATTCAC GAAGATTCAC GAAACCCAG GAAGATTCAC GAAACCCAG GAAGATTCAC GAACTTCTCC CTCTCTCTCCC CTTTCTCCC CTTTCTCC CTTTCTCTC CTTTCTCT CTTTCTCTC CTTTCTCT CTTTCTCT CTTTCTC CTTTCTCT CTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTCT CTTTTTCT CTTTTTCT CTTTTCT CTTTTTT	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTCATGAT AGTTAATAGAC CGATTAATAGAC CGATTATTGA TCATACTGCT AGTTAAGAGC CGATTATTGA TCATACTGCT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTCATACTGT GAGAGGACCA TCAATACTTT AGAACATTTC AGACATCTC TCAGAGCCAT AGATCTTT AGACCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT AGACTCCAT CAGCCATTCAA ATATTTCCAT AGACAATAC CTTCATCAGC CTAGCCATTA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTTCG CTCGGAAATG CCTGGATTGG GTCCTATACA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCAGA CTTATCCATT TGGAGTCGAA GAACCTTCTG GACTACTAC CCAGTTGGCT TCAGTACACT TCAGTACACT TCAGTACCAT CTCATACCAT AAATACAC CCCTGAATTA AGAAGGCGCT ACCCAGATT AAATACAGC CCCTGAATTA GATTACCTAACT GACTAACCT ACCCAGATT GACTAACCT CCTTCAGA AATGATGC ATCCTTAAAT CCTTGATACT CTTGATACT CTTCAGAAGAA ATGATGTC TTCAGAAGAA AGACATAACA CACTGAGATA CACTGAGATA CACTGAGATA	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GCAGCACTCA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTGAG ACTTTTCAA GCAACACTCA GCACACTCA GCACACTCA GCACACTTG GCACACTTG GAAACAATT GGAAACAATT GGAACAGTTG GACCACTGA ATTGTGAACTG ATTGTGAACTG ATTGTGAACTG ATTGTGAACTG ACTGTGACTC CCCAACAA TCCACCACAA TCCACTACCC ACTGTGACTC CTCAAACTG ACAGTTTCTA GCACGTACG CCTTATACCAC CTTATACCAC CTTATACCAC CTTATACCAC CCTTATACCAC CCTTATACCAC CCTCACCAC CCTCTTGACCC CTTATACCAC CCTCACCAC CCTCTTGACCC CCTTATACCAC CCTCACCAC CCTCTTGACT CCACCACCAC CCTCTTGACCC CCTTATACCAC CCTCTTGACCC CCTCTTGACT CCACCACCC CCTCTTGACCC CCTCTTGACT CCACCACCC CCTCTTGACT CCACCACCC CCTCTTGACCC CCTCTTGACT CCACCACCC CCTCTTGACT CCACCACCC CCTCTTGACT CCCTCTTGATACCC CCTCTTGATACCC CCTCTTGATT	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAACAA ATATTGATGA GGGATAAACC ATCTACTAA AGATAACTTT AAGGACAAGTA ATGATTGAGGA GTTTTGAGGA GTTTTGAGGA GTTTTGGGAA ACTGGATTGT AAGTTCTTAC TTCGAGAGCA AGATTCTTAC TTCGAGAGCA AGATTCTTAC TTCTCAGAT TTGTCGACAT TTGTCGACAT TTGTCGACAT AAGAAATAAT CTGGTAGAG AACACTACAA AAGGAAGTGA AACCAGTCAC TTCTTAGATC TAACAGATT AACTGCAC TTCTTAGATC AACTGCAC TTCTTAGATC AACTGCAC ACTCTCTAGAT AATCTTCTAGAT AATCTTCTAGAT AATCTTCTAG AAGGGTATAT AATCTTCTAG AAGGGTATAT AATCTTCTAGAT AATCTTCTAGAT AATCTTCTAGAT AATCTTCTAGATC AACTGCTAG AACTGCTAG AACTGCTAG AACTGCTAG AACTGCTAG AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGA AACTGCTAGAAC AATCTTCTAGATC AATCTTCTAGATC AATCTTCTAGATC	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGGA ATTTCCACTT TCACTGGGGA ATTTCCACTT TTACATTTACA TTACATTTACA TTACATTTACA TTATACATTTACA TTATACATTTACA TTATACATTACA TCTTGTTACA GCCATACGAG AGCAGTTGT CCACAGACTACA ACAGACCATA CCACACTGT CCACACTGT TCACACTTCCTCCACATTCCCCAAATGCCCCCAATTCCCCCAATGCCCCCAATTCCCCCAATGCCCCCAATTCCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCAGAGGAAGAGAGAG	120 180 240 360 420 480 660 720 900 900 1020 1140 1260 1320 1380 1440 1500 1560 1620 1680 1680 1680 1920 2040 2160 2160 2220
50556065707580	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA AACACATTCA GTCAGCGGAG GAAATGCAATA GAGATGCAAAA GGAAAAGGGA TTAGATCCAT AATGCTCAT AATGCTCAT ACAGTTAGCA TCTGGTTATG TCTCTAGAC CAGTTCAGAAC CAGTTCAGAC AGTTCAGAAC CAGTTGATC TGCACTAATG AATCCTGAAC AAGGGGGAA AACCAAATCA AAGGGTGATT AACTTGACC GAAGACTCT AACTTCTCG GAAAACCCAG GAAGATTCAA GGAACTTCTCA GAAACCCAG GAAGATTCAA GTTTGTTCTCAGCTTTCTCTCTCTCTCTCTCTCTCTTCTCTCTC	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TCTACTGCTT AGTTAAGAGC CGATTATGA TCTACTGTT AGTTAAGAGC CGATTATGA TCATACTGTT TGACATCTCC TCTCTGAAAG TCATACTGTT CAGAGACTCT CAGAAAATGT CAGAGACCT TCATATTGA TCATATTGG TGAACTTTT AGACATTTT AGACATTTT AGACATTTT AGACATTTGA TCATTCCT CAGCTCTT CAGCTCTT CAGCTCTT CAGCTCTT CAGCTCTT CAGCTCTT CAGCTCTT CAGCTCTTT CAGCTCTT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCTAT CAGCTCAT CAGCTCTAT CAGCCCAGT	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCCCCTCC CTCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCAGAG TTATCCATT TGGATCAGAG TCCTGCACA CCAGTTGGCT TCAGGCTGAC TTATCACTT TCAGGCTGAC TTATGATACT TCAGGCTGAC CCAGTTGGCT TCAGGCTGAC TATGATACAC TATGATACC AACCAAGCAT AAAATACAGC CCCTGAATTA AGAAGCAGCT GCTACCCAGATT GACTACCAGAT GACTACCAGAT CCCTGAATTA AGAAGGCGC ATCTTTAAAT GCTTCTCAG AAATGATGGC ATCTTTAAAT CACTTCACG AATGATGAC TCATCTCAGA AATGATGAC ATCTTTGATACT TCAGGAAGAA AGACATAACA CACTGAGATA AGACATAACA GATGTCACAG	TCTATACACT CTCTCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAAATTAA AAGCAAGCA CGATTTTCAA TTGTTTGAG AGTGTTAGTC CCAAACTCAA GACACAGTTG GGAAACAATT GGAAAGAAT GGAAACAATT GGAAAGCAA ATGGTTAGTG CCAGAGAATT TGACCACAA ATGGAACTGA ATGGTTAGTG ACCACTACAA ACCACTCACAA TCCCCAACAC ACCACTTCCC ACTGGACTG CCACTTCCC CACTTCCC CACTTCCC CCTTATACCAC TCACACACC CCTTATACCAC TCACACACC CCTTATACCAC GCACACACCC CCTTATACCAC CCTTTGACTG GCACACACCC CCTTTGACTG GCACACACCC CCTTTGACTG CCTTTTGACTG GCACACACCC CCTTTTGACTG GCACCCCCCACA CCTTTTGACTG GCACCCCCCACA CCTTTTGACTG CGTCCCTCCAC	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAACA ATCACACTA AGGATAACAA AGGATAAACA ATCACACTA AGGATAAAC ATCTCACTAA AGGATAACTT AAGGACAAGTA ACTGGACAGGA GTTTTGGGAA CTGACAAGTA ACTGGATTGT AAGTTCTTAC TTCGAGAGCA AGATTCATGA ATTCTTCAGAT TTGTCGACAT TTGTCGACAT TTGTCGACAT AGATGCTA AGATGCTA AGATGCTA AGAGAAGTAA ACCAGTCA AAGAGTACAA AACAGTCAC AACTGCCACC TTCTTAGATC TAACAGATTA ATTCTTCAGG ATCTTCAGA ATCTTCAGAAA TTACAGATCT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGGA ATTTCCACTT TCACTGGGAA AGAAATTTG GCAGGCTGCT TTACATTTAA AGAAATTTG TTTTAAAGAT AATGCAACAA ACAGTACAAG TCATGATTAC TTTGTTACA GCCTACTGAT CAGAGACTTA TCAGAGAGCAG TCACACATATG TCACACTTGGA TCACACTCGG ATTTTCCTCC AAATGCTTCC GGAGGGAAAT AGGCAGAAG GGACAACCAAG GGACAACCAAG GGACAACCAAG	120 180 240 360 420 480 660 720 840 900 900 1080 1140 1260 1380 1440 1560 1560 1680 1860 1980 2040 2100 22100 22100 2280
505560657075	CACACATACG CAAAAAAAAC CGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAACAATTCA GTCAGCGGAG AAATGCAATA GACATTCAA GGAAAAGGGA TTAGATCCAT ACAGTTAGAC TTTCATAGAC TTTCATAGAC TTTCTAGAC TTTCTAGAC TTCTCTAGAC TTCTCTAGAC TTCTCTAGAC TGGCAAATA CAGTTGGATC AGTTCAGAAC AGGTGCTATTC AACTCTGAAC GAAGAGGGAAA AACCAAATC AAGGGTGATG AACTCTGAG GAAAACCCAA GAAGGTACTT AACTTGTCGG AGTTTATTCAGAC GAAGATCAA GCAAATCAA GCAACTTCTG GAAAACCCAA GTTGTTCTCC CCTTTTCTC CCTTTTCTC CCTTTTCTC CCTTTTCTC CATTATTCTA	CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTAATTGA TCTATCTGTT AGTTAAGAC CGATTATGA TCATACTGT TGACATCTC TCTCTGAAAG TCATCTGT TGACATCTC TCTCTGAAAG TCATGCTGAT AGGGTGTTTT CAGAAAATGT CTGAGTCGT TGATATTTA AGACATTAT AGACATTTT AGACATTT AGACATTCC TCCCAATAC TCATCTCT TGATCTCT TGATCTTT AGACCATTT AGACCATT CAGCTCTT AGACCATT AGACCATT AGACCATT AGACCATT AGACCATTAC CTTCATCAGG CTAGCTCTAC AGACTAATTA CTATCCCATT AGACCATTAC AGACTAATTA CTATCCCATT AGACCATTAC AGACTAATTA CTATCCCATT AGACCATTAC AGACTAATTA CAGGCCCCTT CCTTCACCAGG CTAGCTCTAC AGACTAATTA CAGGCCCCAGT CCTTCACCCA AGACTAATTA AGACTAATTA CAGGCCCCAGT CCTTCACCCA AGACTAATTA AGACTATTA	CTCACTTCGA CTCACTTCGA CTCACTTCGA CTCACTTCG CTCGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACA AGAACTTAAA TGGGAAACA AATGGTGTTT TGGATCGAG CTTATCCATT TGGAGTCGAA ACATGTGCT GCCTGCACA CCAGTTGCT TCAGCCACA CTATACACT TCAGCCAAT AAAATACAC CCCTGAATTA AGAAGGCGTT ACCCCAGATTA ACCAACGAT ACCTTCTCG ACCTTAACTA CCCCAGATTA AGAAGGCGTT CCCTGAATTA AGAAGGCGTT CCCTGAATTA CCCTGAATTA AGAAGGCGTT TCTCAGG ATCTTTAAAT GACTTCTCAG ATATGATGC CTTCAACTA CACTGAGAAA AGACATAACA CACTGAGATA AGACATAACA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAGATA CACTGAACT CTCCCAACT CTCCCAACT CTCCCAACT CTCCCAACT	TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GCAAGCACTGA TCTCCTATCA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATATTTCAG TTGTTTGAG AGTGTTAGT CCAAACTCAA GCACAGTTG GTTTTTTGTG CAAAACAATT ATGATTTGAG ATGATTTGAG ATGATTAGA GAATTTTGAG AATGATAT GGAAAGAAT ATGATTAGA ATGATTATG AATGATATG ATTGGAACT ATTGGAACT ATTGGAACT ATTGGAACT CCCAACACA ACCCTACACA ACCCTACACA TCCCCTACACA TCCCCTACACA ACCTTCCC ACTGTGACT CTTAAAACT CCTTAAAACT CCTTAAAACT CCTTAAACT GGAGCTGAG ACCTTCCC CTTATACCAC CTTATACCAC CCTTATACCAC CCTTATACCAC CCTTATACCAC CCTTATACCAC CCTTATACCAC GCACAGCCCC CCTGTTGATG GGACTCCCTCA CGGGCTCACA AACGTGGTAC AACGTGGTAT	GGAGGATTAA TGAGAAGCA ACTACAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGGATAAACTT AAGGACAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTTT AAGGACAAA ATGTTGGGACAG ATTGGGACAGT AAGTTCTTAC TTCGAGAGCA AGATTCATGA ATATTGATGA ATACTACAG ATCTTCAGAT ATCTTCAGAT ATCTTCAGAT ACGACTACA AGATGATA ACGACTACA ACATTCACAC ACTTCTAC ACTTCTAGA ACTGCACA ACATTCACAC ACTTCTAGA ACCAGTCAC ACTTCTAGA ATCTTCAGGA ACCAGTCAC ACTTCTAGA ATTCTTCAGG AAGGGTATAT AATCTGCTAG ATCTTCATAT ATGTTGGATC CTCATGCTTT ACTCGCAGAC	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGGA ATTTCCACTT TCACTGGGGA ATTTCCACTT TTACATTTACA TTACATTTACA TTACATTTACA TTATACATTTACA TTATACATTTACA TTATACATTACA TCTTGTTACA GCCATACGAG AGCAGTTGT CCACAGACTACA ACAGACCATA CCACACTGT CCACACTGT TCACACTTCCTCCACATTCCCCAAATGCCCCCAATTCCCCCAATGCCCCCAATTCCCCCAATGCCCCCAATTCCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCCTCCCCAATGCAGAGGAAGAGAGAG	120 180 240 300 360 420 660 720 780 960 1080 1140 1260 1380 1440 1560 1680 1740 1860 1920 2040 2210 2210 2210 2210 2210 2210 22

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	CARARACCCAA.	CTTABCACCT	TTATCCATTT	TGTTTGAGGT	TGGGACAGAA	GAAAATTTGG	720 780
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15	A MOCONCA TER	CACATCTCCT	CCCTGCACAG	ACACAGTTGA	CTGGATTGTT	TTTAAAGATA	900
	CACTTACCAT	CTCTGAAAGC	CAGTTGGCTG	TTTTTTGTGA	AGTTCTTACA	ATGCAACAAT	960
		CATCCTCATC	CACTACTTAC	AAAACAATTT	TCGAGAGCAA	CAGTACAAGT	1020 1080
20	TCTCTAGACA	GGTGTTTTCC	TCATACACTG CAGGCTGACC	GAAAGGAAGA	TACCAGCCTT	CTTGTTACAT	1140
20	0000000000	かんてみのかんなですか	ጥስጥርስጥስርርል	TGATTGAGAA	GTTTGCAGTT	TIGTACCAGC	1200
	NORTH CONTROL	ACACCACCAA	DTADDAACOA	AATTTTTGAC	AGATGGCTAT	CAAGACTTGG	1260
	ORGON ARROT	CAATAATTTC	CTACCCAATA.	TGAGTTATGT	TCTTCAGATA	GTAGCCATAT	1320 1380
25	GCACTAATGG	CTTATATGGA	AAATACAGCG CCTGAATTAA	ACCAACTGAT	TGTCGACATG	AAGGAGGAGG	1440
25	ATCCTGAACT	TGATCTTTTC	GAAGGCGCTA	TTGTGAATCC	TGGTAGAGAC	AGTGCTACAA	1500
	8 CC8 8 8 TC8C	CADDAAGCAA	CCCCAGATTT	CTACCACAAC	ACACTACAAT	CGCATAGGGA	1560
	OCH B BEREAR	TONACCONAC	አርፕልአርርርልፕ	CCCCAACAAG	AGGAAGTGAA	TTCTCTGGAA	1620
20	AGGGTGATGT	TCCCAATACA	TCTTTAAATT	CCACTTCCCA	ACCAGTCACT	CACACTGTGG	1680 1740
30	CAGAAAAAGA	TATTTCCTTG	ACTTCTCAGA AATGATGGCT	CTARARCTGT	TCTTAGATCT	CCACATATGA	1800
	* ~~~~~~~~~	CACTCCAGAA	ጥርርጥጥልልልጥል	CAGTTTCTAT	AACAGAATAT	GAGGAGGAGA	1860
		CA COMMON CA A C.	ርተጥር ልጥል ርጥር	GAGCTGAAGA	TTCTTCAGGC	TCCAGTCCCG	1920 1980
25	CAACTTCTGC	TATCCCATTC	ATCTCTGAGA TATGATGTCC	ACATATCCCA	AGGGTATATA	AATGCTTCCG	2040
35	33C3TTC33C	ምምር አጥር አርር ጥ	TCAGAAGAAT	CACTAAAGGA	TCCTTCTATG	GAGGGAAATG	2100
	それらいちからして	ጥክርርጥርጥክርክ	CACATAACAG	CACAGCCCGA	TGTTGGATCA	GGCAGAGAGA	2160
	COMPROMOTO	CACTAATTAC	ACTGAGATAC	GTGTTGATGA	ATCTGAGAAG	ACAACCAAGT	2220 2280
40	CCTTTTCTGC	AGGCCCAGTG	ATGTCACAGG TTCCCAACTG	ACCULACT	TACAGATCIG	ACCCCATCCT	2340
40	CONCACANCA	CCATTTCCTC	TCCACGGTCA	ACGTGGTATA	CTCGCAGACA	ACCCAACCGG	2400
	mamacaamca	CCCCAGTAAT	ACTACCCATG	AGTCTCGTAT	TGGTCTAGCT	GAGGGGTTGG	2460
	********	CANCCCACTT	ATACCCCTTG	TGATCGTGTC	AGCCCTGACT	TTTATCTGTC	2520 2580
45	TAGTGGTTCT	TGTGGGTATT	CTCATCTACT	TATCCACACC	TCCAACACCT	ATCTTTCCAA	2640
43	ののなくなべなのなる	TOTOGGAGCA	ATTCCAATAA	AGCACTTTCC	AAAGCATGTT	GCAGATTTAC	2700
	3 TO CO 3 TO CO 3 CO	ጥርርርጥጥጥልርጥ	GAAGAATTTG	AGACACTGAA	AGAGTTTTAC	CAGGAAGIGC	2760
	AGAGCTGTAC	TGTTGACTTA	GGTATTACAG	CAGACAGCTC	CAACCACCCA	GACAACAAGC CTAGCACAGC	2820 2880
50	mmocroca a a a	CCATCCCAAA	CTCDCTCDTT	ATATCAATGC	CAATTATGTT	GATGGCTACA	2940
50	3 C3 C3 CC3 3 3	አ ር ር ጥጥ አ ጥ ስ ጥጥ	CCTCCCCAAC	GCCCACTGAA	. ATCCACAGCI	GAAGATTTCI	3000
	CCACAATCAT	ATCCCAACAT	AATGTGGAAG	TTATTGTCAT	GATAACAAAC	CTCGTGGAGA	3060 3120
	AAGGAAGGAG	AAAATGTGAT	CAGTACTGGC	CTGCCGATGG	TACTGTGAGG	TACGGGAACT	3180
55	MARGRADORO	**********	AACCCCTCCC	' AGAAAGGAAG	ACCCAGTGGA	CGTGTGGTCA	3240
55	. UN UN CENTURA		TGGCCCTGACA	TGGGAGTACC	AGAGIACIC	CIGCCAGIGC	3300
•	ጥር እርርጥጥጥርጥ	CAGAAAGGCA	CCCTATGCCA	AGCGCCATGC	AGTGGGGCCI	GLIGICGICC	3360 3420
,	ACTGCAGTGC	TGGAGTTGGA	AGAACAGGCA	CATATATIGI	BANACACAT	ATGTTGCAGC CGTTCACAAA	3480
60		CCTACAAACT	CACCACCAAT	· ATGTCTTCAI	' TCATGATAC	CIGGIIGAGG	3540
00	CCN TO CTT AC	TABACABACT	GAGGTGCTGG	: ACAGTCATAI	TCATGCCTAT	GITAATGCAC	3600
	ポーク・サース アナーア	ጥርርልሮርልርርል	GGCAAAACAA	AGCTAGAGAA	ACAATTCCAC	CICCIGAGCC	3660 3720
	AGTCAAATAT	ACAGCAGAGI	GACTATTCTG	CAGCCCTAAA	TGGCATTTC	AGGGAAAAGA TCCCTGAGTG	3780
65	0303300030	አ / አ / ጥ አ / ጎ አ ጥ /	י אאַירכררידרירין	' ATATCATGGG	CTATTACCAC	AGCAATGAAT	3840
0.5	ምርአ ጥርአ ጥጥ እር	CCAGCACCC	· CTCCTTCATA	CCATCAAGG	\ TTTCTGGAGG	ATGATATGGG	3900
	N CONTRA NTCC	CONNECTO	ያ ርጉጥልጥርልጥጥር	CTGATGGCCA	AAACATGGC	GAAGATGAAT GTCACTCTTA	3960 4020
	TTGTTTACTG	GCCAAATAAA	GATGAGCCTA CTATCTAATC	TAAATIGIGA DAAAAAAA	ACTTAATAT 1	GACTTTATCT	4080
70	TO A A A COTTA C	' አሮአሮሮአሞር አ ግ	ንሞዮንልጥድንዮልም ሳ	AAGTGAGGC	\ CTTTCAGIG	CCTAAATGGC	4140
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	OMOCON BEDC	CCARCCCCC	ስ ከጥር እጥጥር የ ጥር	· ATGATGAGC	A TGGAGGAGT	ACGGCAGGAA	4260 4320
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	5 mmm/2013 5 C 2	N TOTAL TOTAL	ጥ ሲሚጥልጥጥጥ	T TCTCTATTG	A TTTTAACAG	A AAATTICAAT	4800
	mm = m = c = c c c		~ እአከርሞኒስርኒር	A AAATGTTTG	T TTTTAGTGT	C AAATTTTTAG	4860 4920
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	TTTAGTTTA	A TGACGTAGT	T CATTAGCTG	G TCTTACTCT	A CCAGTTTC	T GACATTGTAT	5160

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						CATTGTTCAA	5280
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15			LPNMSYVLQI				300
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			SLNSTSQPVT				420
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20			YDVLIPESAR TEIRVDESEK				600
20			STVNVVYSQT				660
			LIYWRKCFQT				720
	IPIKHFPKHV	ADLHASSGFT	EEFETLKEFY	QEVQSCTVDL	GITADSSNHP	DNKHKNRYIN	780
25	IVAYDHSRVK	LAQLAEKDGK	LTDYINANYV	DGYNRPKAYI	AAQGPLKSTA	EDFWRMIWEH	840
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	KGSQKGRPSG	RVVTQYHYTQ	WPDMGVPEYS VNIFGFLKHI	LPVLTPVRKA	AYAKRHAVGP	TATEATLOUET	960 1020
	RIGITIVEDS	UNALITECED	GKTKLEKQFQ	LT-SOSMILLAGI	DYSAALKOCN	REKNRTSSII	1080
			NASYIMGYYQ				1140
30			DEPINCESFK				1200
			SKTFELISVI				1260
			NLMRPGVFAD	IEQYQFLYKV	ILSLVSTRQE	enpstsldsn	1320
	GAALPDGNIA	ESLESLV					
35	Sea ID NO.	187 DNA sec	mience				
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			CTCCCCCTCC				120
			TCTGGAAATG				180
			CCTGGATTGG				240
45			GTCCTATACA				300
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			GAAACTTAAA				420
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50			TGGATCAGAG				600
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			GGACTACTTA				1020
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C O	AGTTCAGAAC	CAGAAAATGT	TCAGGCTGAC	CCAGAGAATT	ATACCAGCCT	TCTTGTTACA	1140
60 .	TGGGAAAGAC	CTCGAGTCGT	TTATGATACC	ATGATTGAGA	AGTTTGCAGT	TTTGTACCAG	1200
						TCAAGACTTG	
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75						AAATGCTTCC	
13						GGAGGGAAAT	
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						GGAAATGCCA	
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						TGAGGGGTTG	
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						TGCACACTTT TATCTTTCCA	
85						TGCAGATTTA	
-	CATGCAAGTA	GTGGGTTTAC	TGAAGAATTT	GAGACACTGA	AAGAGTTTTA	CCAGGAAGTG	2760
	CAGAGCTGTA	CTGTTGACTT	AGGTATTACA	GCAGACAGCT	CCAACCACCC	AGACAACAAG	2820

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       AACAGACCAA AAGCTTATAT TGCTGCCCAA GGCCCACTGA AATCCACAGC TGAAGATTTC
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                                                                             4080
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                                                                             4140
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       TARAACACTC TTCCATATGA TATTCAACAT TTTACAACTG CAGTATTCAC CTARAGTAGA AATAATCTGT TACTTATTGT AAATACTGCC CTAGTGTCTC CATGGACCAA ATTTATATTT
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                                                                             5220
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       ATACCTTCAT TTTGAAAGAA GTTTTTATGA GAATAACACC TTACCAAACA TTGTTCAAAT
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       Seq ID NO: 188 Protein sequence:
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       MRILKRFLAC IQLLCVCRLD WANGYYROOR KLVEEIGWSY TGALNQKNWG KKYPTCNSPK
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55
                                                                              240
       TDTVDWIVFK DTVSISESQL AVFCEVLTMQ QSGYVMLMDY LQNNFREQQY KFSRQVFSSY
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       TGKEEIHEAV CSSEPENVQA DPENYTSLLV TWERPRVYYD TMIEKFAVLY QQLDGEDQTK
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       LIGTEEIIKE EEEGKDIEEG AIVNPGRDSA TNOIRKKEPO ISTTTHYNRI GTKYNEAKTN
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                                                                              600
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       LVIVSALTFI CLVVLVGILI YWRKCFQTAH FYLEDSTSPR VISTPPTPIF PISDDVGAIP
                                                                              840
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                                                                              900
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       EVIVMITALV EKGRRKCDQY WPADGSEEYG NFLVTQKSVQ VLAYYTVRNF TLRNTKIKKG
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       SQKGRPSGRV VTQYHYTQWP DMGVPEYSLP VLTFVRKAAY AKRHAVGPVV VHCSAGVGRT
                                                                             1080
70
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                                                                             1200
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80
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                                                                              420
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		GACCTGACGC					720
35		ACCATTGATG					780
		GTCTGTTCCT					840
		TGGTCTGTCA AACAGTGCAC					900 960
		CGCAGTGGGG					1020
40		TCTTTTGAGC					1080
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		GATGATGTGT					1200
		CTGGATGAAA GCATCCCCAA					1260 1320
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,,,		CAGTCTCCCA					1440
		CTGCCTCTGT					1500
		TGCTCAGAGT TGTCCAACAG					1560 1620
50		AACACATTGT					1680
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		CAAAATGTGC					1800
		CCGTTTAAAT					1860
55		TAAAAAGTTT					1920 1980
		TCCCCAGCCT					2040
	CACCAATTCT	GGCTGTCAAT	GGGGAGAAAT	AAACCAACAA	CTTATAATTG	TGACACCAGA	2100
•						AATACTGCAG	
60		AGAGTTTATA				ACTGAGGCAA ACCTGATTCC	2220 2280
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						TATAAAAATT	
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						GTTATGCTTG	
						ATGTTACAGC	
70						ACACAGAAAT	2820
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						TCCTGTGTTG	
						TTTGGAAAGG	3060
75		AAACCAAGCT					3120
75						TGGTGCATGT	3180
٠.						AGACACCCCT GGGGAACCCA	
						GCTACCAAAA	
00	ACATTITCGG	TTCAAGAAAA	GTGAGATGAT	GGTAGTACTG	GTTTCTGGTG	AAATTGAAAA	3420
80						CCATTCAAAA	
						GGTTTNCTTA TAGGGGGTGT	3540 3600
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85						CTTGGGCCTC	
						AATACCAAAT	
	GGAIAIATTT	. CI I'TAGGAT	AMCCI I IGAA	COMMINI	. CARIMACAA	TAGTACATCT	7300

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	WO 02/	080443					
			GTATAAGGAA				3960
			GTCCAAAATT				4020
			TTCAAGGCAA				4080
~			CAATTTTACC				4140
5			ACACACATCA				4200
	ACTTCTCCCT	CTTTTTTACA	CACACACACA	CACACACACA	CACAATCCAT	CTCTTGCTTG	4260
	TTCCTACCTC	CCTGATTTTT	CTTCCCTACA	GAAATAGAAA	TAGGGACAAA	GAAGGGGAAA	4320
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	ATTGAGAGAA	AAGCTCCTTT	TCTCTTCACT	GTTTTGGAAA	GGATAGCCAT	TAGCATGACT	4440
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	GAAGGAACCT	TAAGATCACA	TCATCTACTC	CTCTACTCCA	AATTTCTCAT	TCTTCAGGCC	4560
			TAAAGTAATT				4620
			TCTCCTGGCT				4680
			CTGAGAGTTA				4740
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13			CATCCACATG				4860
			ATGGGGTCCC				4920
			TCTGAATAGC				4980
							5040
20			TCTCCATCAG				
20			GTTTCTTGTT				5100
			TTTCTGAGAT				5160
			TTTNTTTTAG				5220
			GCCCTCATCC				5280
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25	TAAACATTGG	CTCCTGTGTT	TGCACCAAAA	TAGCAAGCTG	TGTGCTCTAT	ACACTCTTCC	5400
	CATCGTCTTG	TGTACACTGC	TCCTGTGGCC	TTCCACAGCA	GAAACCAGGG	CAAAAGGGTC	5460
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	TTCAGTTNTA	AGAGACCTCC	TTCTGGGCTT	ACCCCACTCC	TCAGGTACTT	CTCTCTCCTT	5580
	CCTCCTTCTC	CTCCACAGTC	ACAAGTAACC	AAGGAACCTG	AAAGTGGATG	TGTAGCTATT	5640
30			GAGATTCTTC				5700
•			ACAAAATTTT				5760
			CCTGCACATT				5820
			NAACAGGATG				5880
			ATCACCCCAC				5940
35			AGTGGTTTGA				6000
33							6060
			TTTTTTAAAA				
			CCAGGCCTTA				6120
			CTCCCAAGTG				6180
40			CCATCTATCC				6240
40			AAATTGTTTT				6300
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			GTGACATTGG				6420
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			TCACCTCCCA				6540
45	CGAGTAGCTG	GGATTACAGG	CGTGCGCCAC	CAAATCTGGC	TATTTTTTTA	TATTATTAT	6600
	TTTTAGTAGA	GATGGGGTTT	CACCATGTTG	GCCAGACTGG	TCTTGAACTC	TTGGCCTCAA	6660
	ATTATCTGCC	CACCTCGGCC	TCCCAAAGTG	CTGGGATTAC	AGGCATGAGC	ACCATGCCCA	6720
	GCTGACAAGA	CTAATTTTTT	ATCCCTTGGT	TTATTGGCTT	CAACATCTTC	TGGAATCAGA	6780
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			GTAAGGCAAA				6960
			GAATCAATGG				7020
			GTCTAGTTTC				7080
			AAACATCCAT				7140
55			TTTTTCCTTC				7200
55			CTTGGGACTG				7260
			TCAGAAGTAT				7320
			TTCGTAATCC				7380
<i>c</i> Λ	TATAGCTCAT	GTATCTTTAG	GTCTTTGCCT	TCCAAGCACT	GTACAGAATA	CTTTGTGGTT	7440
60	CCTTTTTAGT	CTGACATTTT	GTGGAGCAGT	GAAGCGTGCT	CAGAGACATA	ATCAGCTGAA	7500
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						AATTTCTCCT	
•	GTCATTTGTT	TTGGCTGGAT	GGGGGTGGGG	GAGTAATTGC	TTAAAGTTTT	ACCATTACAC	7680
							7740
65						TATACTATAA	
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	CTTGACACCA	ACTGTTCATG	ATACTGAATA	GACAGTCCAT	ATAAGAGAAA	TTAGTGGACC	7920
						CTTGGAAATG	
						TTTATAAGTT	
70	ATCCANANCO	GATTTCARCA	ACTANGACCT	TATGCCAAAA	TCTCTCCAAT	GTATGGTCCT	8100
, ,	CTAATATATT	CONCOMMON	CCCV VACAACC	COTTATEACT	TGTATACAAC	TAATGCATGT	8160
	OTHER TRIBLE	THE THE PERSON OF THE PERSON O	CCAUTOUTC	TA DCTCTTGACT	TO THE TANK	GATCACCTTT	8220
	TTTATTGAAT	TITIGCATTIC	CCACGIGIGG	TAAGICIIIA	MAMIGITITI	CATCACCITI	0220
						GAGAAAGTTT	
75						GTACCTGTAG	
13						TGACTAGCTT	
	TGGAGAGTAA	ATGCATGGTA	TTGTACATCA	CATTTCTTAA	CTCGTTTTAA	CCTCTGAAAA	8460
	GAATATATTC	TTCTTTGTAG	TCCTTCTTCC	CACCCCCTTG	CCCTCTCCCT	CTCCCTGCTC	8520
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00						ATCTCTCAGC	
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	TCAGTGGTAA	ATTGGGGTTG	TATTGGCCAT	TGATTACATT	CAGGATTGAA	TAGTTTTCAG	8760
	AATCACATGT	AATCCAAAGA	CAGTAGGTAG	TGATGTCCCT	TATCCCTGCA	GCTGTTTTAA	8820
						AAAAAAAAGA	
	AAAAATGAGA	GAAATAAAAC	AGATATTTAA	GAACTTTAGC	CACCTATTTA	GAATAGTTAT	8940
85	AGCCAGAAAA	AAAAACAAGG	GCATGAGTTC	AAATGCATTA	CTATCAGTGT	CCTAGGCAAT	9000
	ACCEDACCEA	Calculator y y dea	CTCATTCAAA	AGCAGTATTT	CAAGAGGCAT	TCTCCTTTTT	9060
	ACCIVACCIA	ACCCCACATA	GIGHTICHM	MANAGEMENTS.	CCCCCATAA	CCAGCTGGAG	9120
	**************************************	ACCCCACTIG	GWCIGGIWG	1110010400		CIUMU	7120

5	CAGACCCTTT AATGAGGGCT TAAGGTGCTG AGCTGCTCTT	TTCTTGGGTC TGAACTGTGA GCAATTATTT	TGGCTCTCAA	AAGGTTGTCT TTCGCAGCAT GCTCTGTTCT	AGAGAAGTTT TCAATACCAG TCATCGCATT	GCCATGTGTG GCAGCCAAAG CTCATTTCTG	9180 9240 9300 9360
3			GTAATGTCAT	TTTCCAAAAA	TAAAATTTGA	TTTCAAT	
10		214 Protein cession #: 1					
	1	11	21	31	41	51	
15	GDRYFKGIVY EGESYVCSSD LVTIIRSGVK DFFGDDDVFI	AVSSDRFRSF NFFKKVEYTK PRKAVRVLLN ACGPEKFRYA	! SRMNGLPSPT DALLADLTRS NVNPNWSVNV KKTAHSFEQV QDDFSLDENE	LSDNINLPQG KTSANMKAPQ LTDITEAIKL CRVMKGNPSA	VRYIYTIDGS SLASSNSAQA ETGVVKKLYT TAGPKASPTP	RKIGSMDELE RENKDFVRPK LDGKQVTCLH QKTSAKSPGP	60 120 180 240 300
20	MRRSKSPADS	angtsssqls	TPKSKQSPIS	TPTSPGSLRK	HKDLYLPLSL	DDSDSLGDSM	
25	Nucleic Ac	215 DNA sec id Accession lence: 312.	1 #: NM_1304	167			
2,7	1	11	21	31	41	51	
	0001001000))	manamanaa		CA TOTAL COMP	50
30	CTTTCCAACA GTCTTCCTGG TCCTGTGGCA CCCAGGTCGT AAGTGAGAGA	TCTTCGTTCT TAATTTAGTT CAGTCCGTGG GATGCAGGCG TATGAGTGAG	AAGGAGAGGT TTCTCACTGA GTGAGTGAAT CTTTGAGGGA CCATGGGCCG CATGTAACAA GGACCTGTGA	CCGAGACTCA GTGTGGAGGA AAAGGGCCTC GTAATCGTGG GATCCCAATC	GCCGGTAGGT GCCAGCGGGC GCGGTGGTCC CTGGGCTGGA CTCAGAAAGA	CTGCAGAGTG TTAGGACAGG TCCGCCTTCC ACGAGGGAGG GGAAATGACC	60 120 180 240 300 360 420
35	AAGAAGAGGA AAGGAGCACC TTAAGATAGA	ACCACCAACT TGCTGTTCAA GGATGCACCT	GATAATCAGG GGGACTGATG GGAGATGGTC GAAGCAGGTG	GTATTGCACC TGGAAGCTTT CTGATGTCAG	TAGTGGGGAG TCAACAGGAA GGAGGGGACT	ATCAAAAATG CTGGCTCTGC CTGCCCACTT	480 540 600 660
40	ATGAAGACTG	AAACCAAGAA	TATTGTTCTT CTGCAAAAAA	ATGCTGGAAA	TTTGACTGCT		720
45		216 Protein					
	1	11	21	31	41	51	
50			 QPVGPVIVQQ DAPGDGPDVR			 SGEIKNEGAP	60
55	Nucleic Aci	217 DNA sec id Accession mence: 824	#: NM_0014	76.1			
55	1	11	21	31	41	51	
60		 TGTGAGGCAG	TGCTGTGTGG TATGAGTTGG				60 _.
60	GATGAAGTGG GCAGCTGCTC GCTGATAGCC	AACCAGCAAC AGGAGGGAGA AGGAACAGGG	TCCTGAAGTG ACCTGAAGAA GGATGAGGGA TCACCCACAG	GGGGAACCAG GCATCTGCAG ACTGGGTGTG	CAACTCAACG GTCAAGGGCC AGTGTGAAGA	TCAGGATCCT GAAGCCTGAA TGGTCCTGAT	180 240 300 360
65	CAATCACAGT	GTTAAAAGAA	AAATCCAGAG GACACGTTGA AATAAAGCTT	AATGATGCAG	GCTGCTCCTA		420 480
70		218 Protein cession #: N					
			21 PEVIGPMRPE HPQTGCECED				60
75	Seq ID NO: Nucleic Aci	219 DNA seg	ruence #: NM_0014				
80	1	11	21	31	41	51	
85	AGACAGAGAC GCTTCTCGCT	TGAGCGGCCC CCTCCTGCCC	AACCACCAAC GGCACCGCA GCAGCCCGGG ATCTTTGATC	TGCCTGCGCT CCACCTCCAG	CTGGCTGGGC GAGGGAAGTC	TGCTGCCTCT TGTGATTGCA	60 120 180 240
	TCCGCTGCCT	CAACTGCAAT	GACAACACTG AGGGACCGCT	ATGGCATTCA	CTGCGAGAAG	TGCAAGAATG	300 360

			AACTCTGGAC	CONCORCONCO	TA A A C'CA C'CT	CTCACACCAC	420
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	CCAGATGCGA	CCGAIGICIG	AAGTGTGACT	GTGACCCAGC	TGGCATCGCA	GGGCCCTGTG	540
	ACCAGAGACT	GCIAGACICC	AAGCCAGCTG	TTACTGGAGA	ACCOMPAGE	AGGTGTCGAT	600
5	CACCOURTACEA	TANTOTOGAT	GGGGGGAACC	CTGAGGGCTG	TACCCAGTGT	TTCTGCTATG	660
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	TTCATCAGG	TOTTGATGGC	TGGAAGGCTG	TCCAACGAAA	TGGGTCTCCT	GCAAAGCTCC	780
	AATGGTCACA	GCGCCATCAA	GATGTGTTTA	GCTCAGCCCA	ACGACTAGAC	CCTGTCTATT	840
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10	TTCACTACCC	TOTGGACAGA	GGAGGCAGAC	ACCCATCTGC	CCATGATGTG	ATTCTGGAAG	960
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	TCACCAAGAC	TTACACATTC	AGGTTAAATG	AGCATCCAAG	CAATAATTGG	AGCCCCCAGC	1080
	TGAGTTACTT	TGAGTATCGA	AGGTTACTGC	GGAATCTCAC	AGCCCTCCGC	ATCCGAGCTA	1140
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	AGGCCCTGAT	TTCAAAGGCT	CAGGGTGGTG	ATGGAGTAGT	ACCTGATACA	GAGCTGGAAG	2040
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	CAGAAAGTGA	AGCTTCCTTG	GGAAACACTA	ACATTCCTGC	CTCAGACCAC	TACGTGGGGC	2340
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40	TCCTGGATTC	AGTGTCTCGG	CTTCAGGGAG	TCAGTGATCA	GTCCTTTCAG	CIGGAAGAAG	2760
40	CAAAGAGGAT	CAAACAAAAA	GCGGATTCAC	TCTCAACGCT	GGTAACCAGG	CATAIGGAIG	2820
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45	TCCTTAAAAA	CCTCAGAGAG	TTTGACCTGC TACATCAGCC	AGGTGGACAA	ACATOCCACT	GARGETGARG	3060
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	ACCOMOTOR	MAICCAMGAC	GAGGGGCTGG	TCTTACTCCA	GCAGAAGCTT	TCCCGAGCCA	3480
	AGCCICICAG	CARCACCCAA	CTGCGGCCCA	TCATCTCAGA	GCTGGAAGAG	AGGGCACGTC	3540
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	TATCCTCAGG	TCAACTGACC	TGACCCCATT	CCTGATCCCA	TGGCCAGGTG	GTTGTCTTAT	3840
	TOCACCATAC	TOTALCIONCO	CTGATGCTGG	GCAATGAGGC	AGATAGCACT	GGGTGTGAGA	3900
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	ATAGTCAACT	TATTCTTTGA	GTAATGTGAC	TAAAGGAAAA	AACTTTGACT	TTGCCCAGGC	4080
	ስጥርልልል ጥጥር ጥ	TOTTANTOTO	AGAACAGAGT	GCAACCCAGT	CACACTGTGG	CCAGTAAAAT	4140
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	ATCTTATTTT	CTCAATCTCC	TCTCTCTTTC	CTCCACCCAT	AATAAGAGAA	TGTTCCTACT	4620
	CACACTTCAG	CTGGGTCACA	TCCATCCCTC	CATTCATCCT	TCCATCCATC	TTTCCATCCA	4680
	ጥተል ('CT('CAT'	CCATCCTTCC	AACATATATT	TATTGAGTAC	CTACTGTGTG	CCAGGGGCTG	4740
	GTGĠGACAGT	GGTGACATAG	TCTCTGCCCT	CATAGAGTTG	ATTGTCTAGT	GAGGAAGACA	4800
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80	TGGTGCTGCC	TTGCTTCTGT	ATTTCCTTGG	ATTTTCCTGA	AAGTGTTTTT	AAATAAAGAA	5160
	CAATTGTTAG	ATGCC	•				
			n sequence:				
0.5	Protein Ac	cession #:N	P_005553				
85	_				41	C1	
	1	11	21	31	41	51 I	
	1	1	I	1	ı	1	

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MPALWLGCCL CFSLLLPAAR ATSRREVCDC NGKSRQCIFD RELHROTGNG FRCLNCNDNT
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                                                                              120
       HMLTDAGCTQ DQRLLDSKCD CDPAGIAGPC DAGRCVCKPA VTGERCDRCR SGYYNLDGGN
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       SSAQRLDPVY FVAPAKFLGN QQVSYGQSLS FDYRVDRGGR HPSAHDVILE GAGLRITAPL
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                                                                              360
       DNVTLISARP VSGAPAPWVE QCICPVGYKG QFCQDCASGY KRDSARLGPF GTCIPCNCQG
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                                                                              540
10
       CIHNTAGIYC DQCKAGYFGD PLAPNPADKC RACNCNPMGS EPVGCRSDGT CVCKPGFGGP
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       NCEHGAFSCP ACYNOVKIOM DOFMOOLORM EALISKAOGG DGVVPDTELE GRMQQAEQAL
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15
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                                                                              960
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WO 02/086443

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50	GAGCAACCTC CGACCAGAG GCGGGCCCA ACCTGCACC GCTGTTGGGC GCCCAGTGG CGAGGGGCTG TGACTCCTCTG CTTGGAAGAC	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GGAGTGATAG GATGAGGTGCTG	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCGAGTC ATGGATCGGC CGACAACAACC GACAACCACC GGCCACCGTT GATGGCTGTC	CCGGGCGCGG TCCCGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GGCCTTGG GGCATGAAGT ATTGGGGGTG	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT	120 180 240 300 360 420 480 540
50	GAGCAACCTC CGACCAGAG GCGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCAGGGGCTG TGACTCCTTG CATCCTCCTG TCTTGGAAGAC TCTTGCAGGT ATTCTATGAC	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATGTC GAGTGATGTC CTGGATGTCT CTGGATGTCCT CTGGATGTCCT CTGGATGTCCT CTGGATGTCCT CTGGATGTCCC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCGGGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC	CAGCGCCGCC CGAGCAGGC TTGCCACCT CGAGCGAGTC ATGGATCGGC GAGCACCGGG GCAAGCACCGGG GCAAGCACCGTT GATGGCTGTC AGCATGGTAT CAGGTACGAA	CCGGGCGCGG TCCCCGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT CGTGCCTTGA GGCATGAGGT ATTGGGGGTA TTTGGTGAGA TTTGGTCAGA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAGA CTCTCTCAC	120 180 240 300 360 420 480 540 600 660 720
50 55	GAGCAACCTC CGACCCAGA GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTT TGACTCCTTG CATCCTCCTG CTTGGAAGAC TCTTGCAGGT ATTCTATCAGGT ATTCTATCAGGT TGGGTGGGCTT	AGCTTCTAGT CTTCTCCAGC CCTCAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTTC CCTATGACCC GCTGCTTCTC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGGGGGCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTGCCAC TAGTCCATC	CAGCGCCGCC CGAGCAGGGC CGAGCAGTC ATGGATCGGC CGACAACATC GAGCACCGGG GCAAGCACCGT GATGGCTGCT CAGCATGGTAC AGCATGGTAC AGGGAGCACCACGGTAC AGCAGGAAC GGGAGGTACGAA GGGAGGTGCC	CCGGGCGCGG TCCCCGCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GGCATGAAGT ATTGGGGTG GGCAATAGAA TTTGGTCAGG CTACTTTGCT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGGGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCATCAC GTTCCATCAC GTTCCTTCAC	120 180 240 300 360 420 480 540 600 720 780
50	GAGCAACCTC CGACCCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC CCAGTGG CCAGGGGCTG TGACTCCTTC CATCCTCCTT CATCTCAGAG ATTCTATGAC TGGCTGGGCT TGCTGGAAAACA GAAAGACTAC	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTTCTC ACCTCTTACC ACTCTTACC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATCAG TAGTTGCCAC CAGTCAATGC CAGTCAATGC TCTACCTAG GAGCACAATAG CAAGACACAAG GAGCAAAAAG	CAGCGCCGCC CGAGCAGGC TTGCCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGGG GCAAGCAACC GGCCACCGTT AGCATGGCTGTC AGCATGGTAT CAGGTACGA GGGAGGTGCC GCCCTATCCA GAGAAAATCA	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA GGCATGAAGT ATTGGGGGT GGCAATAGAA TTTGGTCAG CTACTTTGCT AAACCTGCAC TGTTGAAACA	ACCCCAACCC ACCTCTCTCC CGCCTTCTGC CGCGTTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTGT TCGTTCAAGA CTCTCTCAC GTTCCTTCCC CTTCCAGCGG AACCGAAAAT	120 180 240 300 360 420 480 540 600 660 720 780 840 900
50 55	GAGCAACCTC CGACCAGAG GCGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCCCCAGTGG TGACTCCTTG CATCCTCCTG CATCCTCCTG TCTTGGAAGAC TCTTGCAGGT ATTCTATGAC TGGCTGGGCT CCGAAAAACA GAAAGACTAC GGACATTGAG	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCAGGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC TCTGCCTTCT CAACACCAAG GAGGCAAAAG TAACATTAGG	CAGCGCCGCC CGAGCAGGC TTGCCACCT CGAGCAGTC ATGGATCGGC GAGCACCGGG GCAAGCAACCAC GGCACCGTT GATGGCTGTC AGCATGGTAT CAGGTACGAA GGGAGGTGCC GCCCTATCCA ACCATAGAAATCA ACCTTAGAAA	CCGGGCGCGG TCCCCGCCTT GCAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT GGCATGAGA GTGACTGAG GGCAATAGAA TTTGGGGTTG TTTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGATAT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGT TCGTTCAACG CTTCCACGG AACCCGAAAAT GTAATCTGAA	120 180 240 300 360 420 480 540 600 600 720 780 840 900 960
50 55 60	GAGCAACCTC CGACCCAGG GCGGGGCCCA ACCTGCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTT CATCCTCCTG CTTGGAAGAC TCTTGCAGGT ATTCTATGAG TGGCTGGGCT CCGAAAAACA GAAAGACTAC GGACATTGAG GTATGGTATT AAACATGGCT	AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT TGGATGTCAT GAGTGATAGA GATGAGGTGC CTGGCTATTT CCTATGACC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT TCAAAACAAA TAATCTTATT	ATCCAGACTC GGCGGCGCG GGAGTCCGGG GCGGGGCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC CAGTCAATGC TCTGCCTTCC CAACACCAAG GAGGCAAAAG TAACATTAGG TAAACAAACA	CAGCGCCGCC CGAGCAGGG TTGCCCACCT CGAGCGAGTC ATGGATCGGC GGACAACATC GAGCACCGTT GATGGCTGTC AGCATGGTAC AGCATGGTACA CGGGGGGGCCCCTATCCA ACCTTAGAAT TCCTCAATAT TCCTCAATAT	CCGGGCGCGG TCCCCGCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT ATTGGGTTGA GGCAATAGAT ATTGGGTTGA CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGGTATT TTTGGTTATT ATTGGGTATT ATTGGGTATT ATTGGGTATT AGGAGGGAAA AGGAGGGAAA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAGTCTT TGGTGGTTGG GTATGAAGT CTCTCTCAC GTTCCTCAC GTTCCTCAC GTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTACCAT	120 180 240 300 360 420 480 540 600 720 780 840 900 960 1020
50 55	GAGCAACCTC CGACCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CCAGCGGCTCC TGACTCCTTC CATCCTCTC ATTCTAGAAGAC TCTTGCAGGT ATTCTATGAC GGAAAAACA GGAAGACTAC GGACATTGAG GTATGGTATT AAACATGGCT TTGTATTACT TTGTATTACT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATTAGT AGAAGATCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGCACATT CAAACACAAG TAACATTAGG CAAACAAACA TTATCTTTT GAGTAATCAT GAGTAATCAT GAGTAATCAT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GACCACCGTT AGCATGGCTGTC AGCATGGTAT CAGGTACGAA GGGAGGTGCC GCCCTATCCA GCCTAGCAA ACCTTAGAAT TCCTCAATAT ACTCAAATGG	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA GGCATGAAGT GTGACTGCA TTTGGGGGT GTTTGATCAG CTACTTTGCT TAAACTTGAACT TTTGGTAAACA TTTGGTAAACA TTTGGTAAACA GGGAAGGGAA	ACCCCAACCC ACCTTCTGC CGCCTTCTGC CGCGTTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTGT TCGTTCAAGA CTCTTCAC GTTCCTTCAC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTACCAT GCTCCTTAAA	120 180 240 300 360 420 480 540 600 720 780 840 900 960 1020
50 55 60	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG CCGAGGGGCTG TGACTCCTG CATCCTCCTG CATCCTCTG CATCCTCAGAGAC TCTTGCAGGC TCTTGCAGGC TCGGAAAACA GAAAGACTAC GGACATTGG GTATGGTATT AAACATGGCT TTGTATTACT TATATATACT TATATATAGT	AGCTTCTAGT CTTCTCCAGC GCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGATAG GATGAGTGATAG CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGGACACA TACAACAAA TAATCTTATT GCTTCCCATT TATGTATATT TAGTATATAGC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTTCCTGGG CCTTCCTGGG CCTTCCTGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGCCTTCT CAACACCAAG GAGGCAAAAG GAGCAAAAG TAACATTAGG CAAACAAACA TTATCTTCTT GAGTAATCAT TACATGTTAAAA	CAGCGCCGCC CGAGCAGGC CGAGCAGTC ATGGATCGGC CGACACATC ATGGATCGGC GCACACCGT GATGGCTGTC GATGGCTGTC GATGGCTGTC GATGGCTGTC GATGGCTGTC AGCATGGTAT AGGATACCA ACCTTAGAAT AAAACCCAT TCCTCAATAT ACTCAATAG TTATTAAAA	CGGGGGGGGGGGTCCCCGCCTTGAACCTCAGTCAGTCAGT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTC TCGTTCAACG CTTCCACG GTTCCACG CTTCCACGGG AACCGAAAAT ACCCAAGA ACTCAGTGCT ATTTACCAT GCTCCTTTAAA ACTCAGTGCT ATTTACAT TGATTAATT	120 180 240 300 360 420 480 540 600 720 780 840 960 1020 1080 1140 1200 1260
50556065	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG CCGAGGGGCTC TGACTCCTG CATCCTCCTG CATCCTCTG CATCCTCAGACAC CCGAAAAACA GGAAAAACA GGACATTGAG GTATGGTATT AAACATGGCT TTGTATTACT TATATATAGA CTCATTATGT CCATATTGAT	AGCTTCTAGT CTTCTCCAGC GCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT TACAAACAAA TAATCTTATT GCTTCCCATT TATGTATATA TGATACTAG GAAGATGTTT	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGTGCGCC CCTTCCTGGG CCTTCCTGGG CCTTCCTGGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC CAGCACATGC CAACACAAG GAGCAAAAG TAACATTAGG CAAACAAACA TTATCTTCTT GAGTAATCAT TACATTAAA ATTGGTATAT	CAGCGCCGCC CGAGCAGGC CGAGCAGTC ATGGATCGGC CGACACATC CGACCACATC CGACCACATC ATGGATCGGC GCACACCGTT GATGGCTGTC GACACGGG GCCACCGTT CAGGTACGAA AGGAGAAATCA ACCTTAGAAT AAAACCCAT TCCTCAATAT ACTCAAATGG TCTATTAAAA TTTCTTTTTC	CGGGGGGGGGGGTCCCCGCCTTGAACCTCCAGATCCAGTGACGCCCCCAGATCCAGTAAGAATTGGGGTAAACTTTGCTAAACCTGCACTGGAAACCAGGGAAGGGGAAGGGGTAATAGACAGAAAAAAAA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTC TCGTTCAACG CTCCTTCAC GTTCCTTCAC GTTCCTGCC CTTCCAGCGG AACCGAAAAT ACCCATGAA ACTCAGTGCT ATTTACCAT GCTCCTTTAAA GATATCTAAT GATATCTAAT ACATATGTAA	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020 1080 1140 1200
50 55 60	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CCAGCGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCAGAGAG ATTCTATGAC TGGCTGGGCT TGGAGAAACA GGACATTGAG GTATGGTATT TATATATAGA CTCATTATTACT CATCATTATTAC CATCATATTAC CATCATATTAC CATCATATTAC CTATATTAC CTATATT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TCATTCTCCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGTGC CCTGGCTATT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAACAAA TAATCTTATT GCTTCCCATT TATGTATATA TGATACTAGC GAAGATGTTT TCATTTACTC AAGGATGAAT	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTTCCTGGG CCAGCACATT CAATCTTTGT AGAAGATTAGCAG CAGCACATT CAATCATTTC CAATCATAGCCAC CAGTCAATGC CAGCACATT CAATCATTT TATACTTCT GAGTAATCAT TACATGTTTT ATACTTTAAAA ATTGGTAATA TTCTTCAATT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGTC ATGGATCGGC GAGCACACTC GAGCACCGTT GATGCATCC GAGCACCGTT CAGGTACCAC GCCACCGTT CAGGTACCAA ACCATACCAA ACCATACCAA ACCATACAA ACCATAGAAT AAAAACCCAT TCCTCAAAAT TCCTCAAAAT TTCTTTTTCT CCTTTTTGTTTTTC CCTTTTTGGGTGTG CTTCATGGGTG CTTCATGGTG CTTCATGGTG CTTCATGGTG CTTCATGGTG CTTCATGGTG CTTCATGGTG CTTCATGGCTT	CCGGGCGCGG CCAGATCCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT ATTGGGGGT ATTGGGGGT AAACTGCAG TGTTGAAACA TTTGGTAAACA TTTGGGTATT GTGTAAAAT AGGAGGAAG ATAGGTAAT ATAGGTAAT ATAGGTAAT CCTTTACAC CCCTTTTCACC GCCCTTTCA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTTCAC GTTCCTCTCAC GTTCCTCCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT ATTTTACCAT GCTCCTTCAA AAATACTATT GCTCCTTAAA AAATACTATT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT ACATATGTAAT AAAGACCTAGC TATACTTATT	120 180 240 360 420 480 540 600 720 780 960 1020 1140 1200 1320 1380 1440
50556065	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG CCGAGGGGCTG TGACTCCTG CATCCTCCTG CTTGGAAGAC TCTTGCAGGT ATTCTATGAC GGACATTAG GGACATTAG GGACATTAG GTATGGTATT AAACATGGCT TATATATACT TATATATAGAC CTCATATTAGT CCATATTACT TATATTATCT TATATTATCT TATATTATCT TATATTATCT TATATTTCT TTTATTTCT TTTATTTCT TTTTTTTCT TTTTTTTCT TTTTTTTT	AGCTTCTAGT CTTCTCCAGC GCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCCTGA GAGTGATAG GATGAGTGC CTGATCTCA GATGAGCCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGTGCGCG CCTTCCTGGG CCTTCCTGGG CCTTCCTGGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAC CAGTCAATCCCTTCC CAACACCAAG GAGCAAAAG GAGCAAAAG AGACATAAGC TATCTTCTT GAGTAATCAT TACATTAAAA ATTGGTATAT TTCTTCATTA TCTTCAATT TCTTCAATT TATACTTA	CAGCGCCGCC CGAGCAGGC CGAGCAGGT CGAGCACATC CGAGCACATC CGAGCACACTC CGAGCACACTC GACACACTC GACACACTC GACACACGG GCAACACTC GACACGGT GATGCATGT AGCATGGTAT CAGGTACGAA ACCTTAGAAT ACATTACAATGA TCCTCAATAT ACTCAATAT TCCTTTTTC GCTTTGGGTG CTTCATGGTT CATCGTATT ACATTTCATAT ACATTTCATAT ACATTTCATATA	CGGGGGGGGG TCCCCGCTT GCAAACTCTC ATGGCAACG GCATCGTCA GTGACCGCC CGTGCCTTGA GGCATCAGT ATTGGGGTA ATTGGGGTA ATTTGGTCAGG CTACTTTGCT AAACCTGCAC TTTGAACGCC TGTTGAAACT ATTGGGTATT ATAGGGGAAG GGGAAGGGT ATAGACAGT ATAGCTATAT CCTTTGCAC GCCCTTTTCA AAGCCCTTATT GCCTACATTT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTGCCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTC TCGTTCAACG CTTCCACGGG AACCCAAAGA ACTCCTTCAC GTTCCTGCC CTTCCAGCGG AACCGAAAAT ACTCAGTGCT ATTTACCAT GCTCCTTAAA ACTCAGTGCT ATTTAATT ACATATGTAA AAGACCTAGC TATACTTATT TGTTTTTTTTT TGTTTTTTTTTT	120 180 240 300 360 420 480 540 600 660 720 780 840 900 960 1020 1140 1200 1260 1320 1380
50556065	GAGCAACCTC CGACCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CCAGCGGCTC TGACTCCTTC CATCCTCTC CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC GGAAAAACA GGAATTGAC TTGTATTACT TATATATACA TCATTATGAT CCATATTGAT CCATATTGAT CAGTCAAATA CTAATTTACC TTATTTTTA TTTCATTGGT AGCCAAGAAG AGCCAAGAAGA GAGCAAGAAG CTAATTGAT CAGTCAAATA CTAATTTACC TTATTTTTA TTCATTGGT AGCCAAAGAAG	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGTGTC CCTGAGTATT CCTATGACCC GCTGCTTTCTC ACCTCTTACC ATACTATCAT ACAAAACAAA TAATCTTATT TGGTTCCCATT TATGTATATA TGATACTAGC GAAGATGTTT TCATTTACTC AAGGATGATT CCATAAATCTT CATATATTC CATATATTC CATATATTC CATATATTC CATATATTC CATATATTC CATATATTC CATATATTC CATATATTATC CATATATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTAC CATATATTATTATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTATTATTAC CATATATTATTAC CATATATTAC CATATATTATTAC CATATATTAC CATATATTATTAC CATATATTAC CATATATTAC CATATATTATTAC CATATATTAC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG CCTATCCCGG GCAGCACATT CAATCTTTGT AGAAGATCAG TAGTTGCCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC TAACACCAAG TAACATCAG TAACATTATG TACATCATT ATACATCATT ATTCTTTAATA TTCTTCAATT ATAGCACTAG TGAATCAGAA CAAACAACA AATCAGAAC TACTTCATTA TCTTTCAATT ATAGCACTAG TGAATCTACAT ATAGCACTAG TAATCAGAAC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGTC ATGGATCGGC CGACACATC GAGCACCGGG GCAAGCAACC GGCCACCGTT GATGGCTGTC ATGGCTACCAA ACATGGTACCAA ACCTAGCAA ACCTAGAAT TCCTAAATGC TCTAAATAG TATCTCTAAA TATCTCTAAA TATCTCTAAA TATCTCTAAA TATCTCTAAA TATCTCTAGCT CATTGGGTG CATCGTTATT TCATATCTCATAT TCATTTTCGGTG CATTCATTAT TCATATT TCATATT TACATTATAT TCATATTTCATAT TTTTGGGTG CATTCATTAT TTTTGGGGCA	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT GTGCCTTGA GCCATCAGT GTGATCAGT CTTTGGTCAG TTTGGTCAG TTTGGTAAACT TTTGGTAAACT GTGTTAAAACT GTGTTAAACT GTGTAAACT TTTGGGAGG ATAGGAAGGGT ATAGGTAAT CCTTTGCCAC GCCCTTTTCA AAGCCCTTAT AATCTTTCTG	ACCCCAACCC ACCTCTCTCC CGCCTTCTGC CGCGTTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGAT CTCTTCAAGA CTCTCTTCAC GTTCCTGCC CTTCCAGCAG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ACTCTTACAC GTTCCTTCAC GTTCCTTCAC GTTCCTGTCC TTCAACAAA AATACTATT GTATTTACTT TGATTTAACT ACATATGTAA AAGACCTAAC TAGTTTTTTT TTGTTTTTGTG TAGTTTTATT TTGTTTTTTTT	120 180 240 360 420 480 540 600 660 720 780 840 900 960 1020 1140 1260 1320 1380 1440 1560 1660 1620
50556065	GAGCAACCTC CGACCAGAG GCGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGGTG TGACTCCTTG CATCCTCTG CATCCTCTG TCTTGGAAGAC TCTTGCAGGT ATTCTATGAC GGACATTGAG GTATGGTATT AAACATGGCT TTGATTACT TTATTATAC CCATTATGAC CTATTATGAC CTATTATGAT CATTATATAC TTATTATAC TTATTATAC TTATTATAC TTATTATAC TTATTATAC TTATTATAC TTATTATAC TTATTATAC TTATTTTTA TTTCATTGGT AGCCAAGAAG GTGATAAATT TTTCATTGGT TTGCTTTGA	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG TCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGC CTGGCTATTA CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGGGCGCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCAC CAGTCAATGCCAC CAGTCAATGC CAACACAAA TAACATTAGG CAAACAAAAA TTACTTCATT ATACTTAAAA ATTGGTAATT TTCTTCATAT TTCTTCATAT TTAGACTTAC TAGACCACT TGAATCAACA CAATCAACA CAATCAACA CAATCAACA CAATCAACA CAATCAACA CAATCAACA CAATCAACA CAATCAACA CCAATTGAGT CCCACACA CCCACTACGCACA CCCACTCCGCACA CCCACTACGCACC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCCACTACCACAC CCACTTCCACAC CCACTTCACT CCACCAC CCACTTCCACAC CCACTTCCACTTCCACAC CCACTT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGTC ATGGATCGGC GGCAACACATC GGCCACCGTT GATGGCTGTC GAGCACCGTT GATGGCTGTC GAGCACCGTT GATGGCTGTC GCCCTATCCA GAGAAAATCA ACCTTAGAAT ACTCTAAAAT TCTCTAAAAT TCTTTTTGCT CTTCATGCGT CATCTTCATGCT CATCTTCATGCT CATCTTCATGCT CATCTCATGCT CATCTTTTTCGCTTCATGCT CATCTTATATAT TTTTGTTTTCATGCT CATCTTATATAT TTTTGATGCT ACATTTCATAT TTTTGAGGCA ACCCCTGTAC AGCTGCATGC	CCGGGCGCGG CCACACG CCCCT CCACACG CCCCT CCCCC CCCCC CCCCC CCCCCC CCCCCC CCCCCC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGGGGCTGCA AGGCCATGTA AGGCCATGTT TGGTGGTTGG GTATGAAGT TTCGTTCAAGA CTCCTTCAC GTTCCTGTCC CTTCCACGGAAAT ACTCAGTGCT ATTTTACCAT ATTTTACCAT GCTCCTTAAA AAATACTATT GTATTTAATT ACATATGTAA AAGACCTAGC TAGGTTCTTAA CATGACTAAT TTGTTTTGTG TAGTTTCTAA CATGACCTAA AGCACTCTTG GGTGTTTTAA	120 180 240 300 360 420 480 540 600 720 780 840 900 1020 1140 1260 1320 1320 1440 1500 1560 1680 1740
5055606570	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTG TGACTCCTCTG CATCCTCCTG CATCTCTGAAGAC TCTTGCAGGT ATTCTATGAC GGACAATACA GGACATTGAC TTGTATTACT TATATATAGA CTCATTATGT CCATATTGAT CCATATTGAT CTATTTTTTT AGCAAGACA TTTTTTTTTC TTTATTTTCT TATATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTTTTTT	AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG CCTGAGCCAG CCTGAGCCAG CCTGAGCCAG AGGATTTACT CGAATCTCA GCAGTGATCTCA GCAGTGATCTC CCTGACTCTAC CCTGCTTTCC CCTGCTTCTC ACCTCTTACC ACTCTACAT CCAAAACAAA ATAATCTAT TGATACTAT TGATACTAG CGAGAGTGTT TCATTACT AAGGATGAT TCATTACT AAGGATGAT TCATTACT CATAATCTT TCTCATCTC AATTTATTAC CCTGTTGACC AAATATTTT CCTTTCC AATTTATTAC CCTGTTGACC AAATATTTTTT TTGATTGACT CAAAATTTTTTTT TTTTATTTGATTTTAC CCTGTTTGACC AAATATTTTTTTTTT	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATGCCGG CCTATGCCGG CCAGCACATT CAATCTTTGT AGAGATGAG AGAGCACAT CAACCAAG CAACAACAACA CAACAACAAC TAACATCTTT TACATGTTT ATACTTAAA ATTCTTCATA ATCTTCATTA ATACTTAAA TTCTTCATTA TACATGTATA TACATGATA TACATGATA TACATGATA TACATCAAC CAACAACAAC CTACACACACACACA	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGTC ATGGATCGGC CGACACATC GGACACACCGG GCAAGCAACC GGCCACCGTT AGGATGGTAT CAGGTACGAA CGGAGAACC GCCCTATCCA GCCTATCCA ACCTTAGAAT ACACTCAAATG TCTCAAATG TCTTATTAAA TTTCTTTTC CTTTGGGTG CATCGTTATT TTTGGAGGCA ATCCCTTGATA TTTCATAT TTTGGAGGCA ATCCCTTGATA CAGTTCATAT ACATTTCATA TTTGGAGGCA ATCCCTTGATA ACACTGTTAT CTTTGGAGCA ATCCCTTGATA CTTGGAGCA ATCCCTTGATA CAGTTGCAT CTTTGGAGCCA ATCCCTTGATA ACTCCTTATCATA CTTGGAGCCA ATCCCTTGATA ACGTTGATGC CTTTATTCATA	CGGGGGGGGG TCCCCGCTT TCCCCGCTT TCCCCGCTT ATGGCAACG GCATCGTCA GTGACCGCC CGTGCCTTGA GGCATCAGT ATTGGGGTA ATTGGGGTAGA TTTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGGTATT GTGTAAAAT ATGGGAAGGGT ATAGGAAGA ATAGGTAAAT ATAGGTAAT CCTTTTCCCAC GCCTTTTTC AACCCTTAT GCCTACATTT AATCTTTCTG TCTGACCCAT TCTGACCCAT GTTTTCACAC GTTTTTATAT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTTCTGC CGCGTGCAA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGT CTCTTCAC GTTCCTTCAC GTTCCTGTCC CTTCCAGGG AACCCAAAAA ACTCAGTGCT ATTTACCAT GCTCCTTCAA GCTCCTTAAA AAATACTATT GTATTTAATT ACATATGTAA AAGACCTAGC TATGTTTTTTTTTT	120 180 240 360 420 480 540 600 660 720 840 900 1020 1140 1200 1380 1440 1500 1560 1620 1680 1740 1800
5055606570	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG CCAGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCTCAGAGAG ATTCTATGAC TGGCTGGGCT ATTCTATGAC TGGAGAACAA GAAAGACTAC GGACATTATGT TATATATAGA CTCATATTGAT CAGTCAAATTA TTTCATTGAT AGCCAAGAAG TTATTTTTA TTTCATTGAT AGCCAAGAAG GTGATAAATT TTTCATTGAT AGCCAAGAAG CTAATTTAC TTTTCATTGAT AGCCAAGAAG CTAATTATATAC TTTCATTGAT AGCCAAGAAG TGATAAATT TTTCATTTGAT CACTAATTTAC TTTCATTTGAT AGCCAAGAAG CTCAATTTAA ACCTTTTTTTA ACCTTTTTTTTA ACCTTTTTTTT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TCATTCTCCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGTGC CTGGCTATT CCTATGACCC GCTGCTTCTC ACCTCTTACC ATACTATAT TGATACTAT TAATCTTATT TCATTATAT TCATTATAT CCATATATCT AAGGATGATT CCCTATTTCCCAATTTATTAT CCTTATCTC AATTATTAT CCCTGTTGACC AATTATTAT CCCTGTTGACC AATATTTATT TCATTTATTAT CCTGTTGACC AATATTTATT TCATTTATTAT CCTGTTGACC AATATTTGT TTGATTGATT TCACCCATTC TTGATTGATT TCACTTCC TTGATTGACT TTGATTGATT TCCCCATTCC TTAATAAGGTG	ATCCAGACTC GGCGGCGCCAG GGAGTCCGGG CCTTCCTGGG CCTTCCTGGG CCAGCACATT CAATCTTTGT AGAAGATGAC TAGTTGCAC CAGTCAATGC CAGTCAATGC CAACACAAG GAGGCAAAG TAACTTTCT TAACATCAAA TAACTTTCT TAACATTACT TAACATCATT TTCTTCAATT TTCTTCAATT ATAGGAACATT ATAGGAACATT ATAGGAACTT TTCTAATT TTCTTCAATT ATAGGACTTAAT TTCTTCAATT ATAGGACC CAAATCAACA CAAATCAACA CCAATTGAGT TTTTAAGGTA TTTTAAGTTA TTGTTTAATT TTCAATT TTCATT TTCAATT TTCATT TTCAATT TTCATT TTCAATT TTCATT TTCATT TTCATT TTCAATT TTCATT	CAGCGCCGCC CGAGCAGGC CTTGCCCACCT CGAGCAGGC CGACCACTC CGAGCAGCT CGAGCACACTC GAGCACCGTG GACACCGTT GATGCCTGTC GAGCACCGTT CAGGTACCAA GGCAGCGTACCACACCGTT CAGGTACCAA CCCTTATCCAA TCCTCAAATG TCTTATAAA TTTCTTTTC CCTTTAGGCT CATCGTTAT TCATCATCAT TTTGGAGGC CTTCATGCT ACATTCATAT TTTGGAGGCA ACCCTGTAC ACCTGCACACA ACCCTGTAC ACCTGCACACA ACCCTGTAC ACCTGCACACA ACCCTGTAC ACCTGTACCACACACACACACACACACACACACACACACA	CCGGGCGCGG TCCCCACACG TCCCCACACG GCCATCGCCACGCC CAGATCCACG GCATCGCCC CAGATCCACG GCCATCGCCC CAGATCCACG GCCATCGCC CAGATCCACG GCCATCGCC CAGATCCACG GCCATCGCC CAGATCCACG GCCATCGC TTTGGCACG TTTTGCTCAC TTTTGAAACA TTTGGTAAAAT AGGAGGAAG ATAGGCATA ATAGGTAAAT CTCCTTATAC CCCTTTTCAC GCCCTTTTCA AACCCTTAT GCCCACTTTC TCTGACCCAT TCTGACCCAT TCTGACCCAC TTTTCCCCCA GTTTTCCCCCA GTTTTTAATATC AGTGCTAGAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTTCT TGGTGGTTGG GTATGAAGT CTCTTCAACAGA CTCTCTTCAC GTTCCTCCC CTTCCAGCCG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT GCTCCTTAAA AAATACTAAT TAATTTACAT TGATTTAATT ACATATGTAAT TGATTTCTAT TAGTTTCTAG CAGCCAAA AGCACTCTG GGTGTTGTAA CCCCTAAACT CCCTAAACT TTCTTGGAGT	120 180 240 360 420 480 540 600 660 720 780 840 900 960 1020 1140 1200 1320 1380 1440 1500 1560 1620 1680 1740 1800 1920
505560657075	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCCCCAGTGG CGAGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCTCATGAGAC TCTTGCAGGT ATTCTATGAC GGACATTGAG GTATGGTATT AAACATGGCT TTGATATACT TATATATACA CCTATATTGAT CCATTATTGAT CAGTCAAATTA TTTCATTGGT AGCCAAGAAG GTGATAAATT TTTCATTGGT AGCCAAGAAG GTGATAAATT TTTCATTGT TATCTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTC CGATAATCTTCC GGATAATCTGC	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG CCTGAGCCAG TCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGC CCTGAGTATAG CCTGAGTATAG CCTATTAGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATATAT TCATTACTAT TCATTACTAT CAAGATGATT TCATTACTC AAGGATGATT CCATATTACTC CAAGATGATT CCTATTCCCATT CTCATTCCCATT CTCATTCCCATT CTCATTCCCATT CTCATTCCCATT CTCATTCCCATT CTCATTCCCATT CTCATTCCC AATATTATTAC CCTGTTGACC AAATATTTGT TTGATTGACT TTGATTGACT TTGATTGACT TTGATTGACT TTGATTGACT TTGATTGATT TCCCCATTCC TAATAAGGTG TTGATTGATT TCAATAGGTG TTGATTGATTT TCAATAGGTG TTGATTGATTT TCAATAGGTG TTGAATAATATT TCAAAATATT TCAAAATATT TCAAAAATAT TCAAAAATAT TCAACAAAATAT TCACCAAAATAT TCACCAAATAT TCACCAAAATAT TCACCAAATAT	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGGGCGCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCAC CAGTCAATGC TCTGCCTTC CAACACAAG GAGGCAAAAG TTACTTCTT GAGTAATCAT TACATTAAAA ATTGGTATT TCTCTCATTA ATAGGACTTC TGAATCACTTC TGAATCACTTC TGAATCACTTC TGAATCACTTC TGAATCACTTC TGAATCACTTC TGAATCACTTC TGAATCTACTT TCTCCTGATT TTACTTACATT TCTCCACACA CCAATTGAGT TTTTAATTGATT TTTTAATTGATT TTTTAATTGATT TGGTCTGTTT TCTCTCTGTA	CAGCGCCGCC CGAGCAGGC CTGACCACT CGAGCAGGC CGACACACT CGAGCACACT CGAGCACACT CGAGCACACT CGAGCACCGT GAGCACCGT GATGCATCGAC GCCACCGTT CAGGTACGAA GGAGGAGTCC GCCCTATCCA GAGAAAATCA AACACTAGAAT ACTCTAAAAT TCTCTAAAAT TCTTTTTC CCTTCATGCT CATCTTAGAGT ACATTTCATAT TTTTGAGGCA ATCCCTGTAC ATCCTGATAT TTTTGAGGCA ATCCCTGTAC ATCCTGATAT TTTTGAGGCA ATCCCTGTAC ATCCTGTAC ATCTTATATA TTTTTTCATA TTTTGAGGCA ATCCCTGTAC ACTCTGATAC ACTCTGATAC ACTCTGATAC ACTCTGATAC ACTCTGATAC ACTCTGATACA ACTCTTATACAT TGTTTTCCCA ACTCTGAACAA GCTGTAACAA GCTGTAACAA	CCGGGCGCGG CCACACCA TCACCACCA TCACCACCA TCACCACCA CCACACCACCA CCACACCACCA CCACACCAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGTTCTGT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTTCAC GTTCCTGCC CTTCCACGGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT GCTCCTTAAA AAATACTATT GTATTTAATT ACATATGTAA AAGACCTAGC TAGACTAAC TAGACTAAG AGCACTCTTG GGTGTTCTAA CCCTAAACT TCATGCCTTAA CCCTAAACT TCATGCCTTAA CCCTAAACT TCATGCCTTAA CCCTAAACT TCATGCGTT TCTCTGGGT TCTTCTGCGTT TCTTCTGCGTT TCTTCTGCGTT TCTTCTCACC	120 180 240 300 360 420 480 600 -660 720 780 840 900 1020 1140 1260 1320 1320 1440 1500 1680 1740 1860 1740 1890
5055606570	GAGCAACCTC CGACCCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCCCCAGTGG CCAGCGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCAGGG ATTCTATGAC CGAAAAACA GGACATTAGT TATATATAGA CTCATTATGT CCATATTAGT CAGTCAAATA CTAATTTAT TTTCATTGGT TATATTTTA TTTCATTGGT AGCCAAGAAG GTGATAAATT TTTGCTTTGA CACATCTTTT TATATCTTCATTGGT CACAACTTTTA ACCTTTTTTT TATATCTTCATTGGT TATATCTTCC GATAATCTCC GATAATCTGT TATATCTTCC GATAATCTGT TATATCTTCC GATAATTTAATT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG GCACCTTCG CCTGAGCCAG GCACCTTCG CCTGAGCCAG GAGTGTTCC CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCCGGG CCTATCCCGG GCAGCACATT CAATCTTTGT AGAACATT CAATCTTCATC TCTGCCTCTC CAGCACATT CAATCTTCT TAATCTTCT TAACACCAAG TAACTTTCT TACATCTTT ATACTTAAAA ATTGCTATA TTCTTCATTA TCTTCATTA TCTCCACACA CCAATTGAGT TTTAAGCTT TCCCTCTTT TCCCTCTTT TCTCTCTTTATACTT TTCTCTCTTTATACTT TTCTCTCTTTATACTTT TCTCTCTTTT TCTCTCTTTT TCTCTCTTTT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGTC ATGGATCGGC CGACACATC GAGCACCGTG GACACCGTG GACACCGTT GATGGCTGTC AGGTACGAA GGCAGGTAT CAGGTACGA GGCAGGTACCA ACCTTAGAAT TCCTTAGAAT TCCTCAAATGG TCTATTATA ACTCAAATGGT CATTCGTTT CACTTTGGGTG CATTCGTTAT ACATTTCATT ACATTTCATTT	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GATGACGCCC CAGATCCAGT ATTGGGGGT ATTGGGGGT ATTGGTCAGA TTTGGTCAGA TTTGGTAAACA TTTGGTAAAAT AGGGAAGGGT ATAGGAGGTAA GGGAAGGGGT ATAGCTCTACAT ATCTTTTCAAACA TTTGCTTAAAAT AGGAAGGGTA TTGGCTATT GCTTATTAAAT ACTCTTATT CCTTTTGCCAC GCCTTTTCA AACCTTATT AATCTTTCTG TCTGACCCAT TCTGACCCAT TCTTACACA GTTTTATATA AGTGCTAGAC AGTTTAGAAC AGTTAGAAC AGTTAGAAC AACTTAGAAC AACTTAGAAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTTCTC CGTCGTTCTCAGGA CTCTCTCAGCG CTTCCAGCGG AACCGAAAAT GTAATCTGA ACTCAGTGCT ATTTTACCT CTCTCTAAA AAATACTATT CTCTTTAATT ACATATTTAATT ACAGCACAAA AGGACCTAGC TATACTTAT TTGTTTTCTG TAGTTTCTAA CCCCTAAACCT CTTCTCAGCGT TTCTTCTG CTTCTTCAC CTTCTTCTG CTCTTTCTTCTG CTCTTTCTTCTG CTCTTTCTT	120 180 240 360 420 480 540 600 660 720 840 900 960 1020 1140 1220 1380 1440 1560 1560 1620 1680 1740 1860 1920 1980 1980 1980
505560657075	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CCAGCGGGTG CATCCTCTG CATCCTCTG CATCCTCTG CATCCTCTG CATCCTCTGCAGGT ATTCTATGAC TGGCTGGGCT TGGAGAACA GGACATTGAG GTATGGTATT TATATATAGA CTCATATTGAT CAGTCAAATTAA CTCATATTTTT TTTCATTGGT AACCAGAGAG GTGATAAATTAA CTTATTTTTT ATTCATTTGT TATATTTTTT ACCATCATTTGAT ACCACACTTTA ACCTTTTTTG TATATCTTCC GATAATCTG TATATTACT TATATTTTTT TATATTTTTT TTTATTTTCT TATATTTTTT TTTATTTTTTT TTTTTTTT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG CCTGAGCCAG CTCATTCTCCG AGGATTTACT TGGATGTCTGA GAGTGATCTGA GAGTGATTGC CCTGAGTCTTC CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCCAG GGAGTCCGGG CCTTCCTGGG CCAGCACATT CAATCTTTGT AGAAGATAGA CAGTCAATGCCAC CAGTCAATGCCAC CAGTCAATGCCAC AGACACATT CAAACACCAAG GAGGCAAAAG ATACTTTCT TACATTATT TACATTATT TACATTATAT TTCTTCAATT ATAGGACTTC TAAATCAAC CAAATCAACA CAATCAGAT TTCTTCAATT ATAGCACTTC TAAATCAGAC CAAATCAGAC CAAATCAGAC TTACATGTTT TTCTTCAATT TTCTTCAATT TTCTTCAATT TTCAATTAGGAC TTAAATGGAAC TTAAATGGTAT TTCATCACTT TTCAATTTAAGCAC TTCACACAC CCAATTGAGT TTTAAATGTAT TTGTTCTTCTTT TCTCTCTGTA TTGAGATAAT ACTCTCATTA AGGACACTGAA	CAGCGCCGCC CGAGCAGGGC CTTGCCCACCT CGAGCAGGTC ATGGATCGGC GAGCACACTC GAGCACCGTT GATGCATCC GAGCACCGTT GATGCTGTC GAGCACCGTT CAGGTACCAA ACCATGGTAT CAGGTACCAA ACCATACCAA ACCATACAA ACCATACAAA TCCTCAAAATCA TCTCTAAAAT TCTCTATACA TTTCTTTTTC CCTTCATGCGT CATCGTTACTAAA TTTCTTTATAAA TTTCTTATAT TTGGAGGCA ACCCTGTACCA ACCTTAGACT CTTATCATAT TTTGGAGGCA ACCCTGTAC ACCTGTACCA ACCTGTACCA ACCTGTACCA ACCTGTACCA CCTTTACCAC GCTTTACCAC GCTTTAACAC GCTGTAACCA GCAGCACCTGC CTTTGAACCA GCAGCACCAC CTTTGAACCA GCAGCACCAC CACACCAC CACACCAC CACACCAC CTTTGAACCA CACACCAC CACACCAC CACACCAC CACACCAC	CCGGGCGCGG TCCCCGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT ATTGGGCGTG TTTGGTCAGA TTTGGTCAGA TTTGGTCAGA TTTGGTAAAA TTTGGTATT GTGTAAAAT AGGAGGAAG ACTTACAT AATCTTTCTA AACCTTTACA GCCCTTTTCA AAGCCCTAT TCTTTCACC AAGCCTTAT TCTTACACT TCTTACACT TCTTACACT AATCTTTCTC AAGCCCTTAT AATCTTTCTC AATCTTTCCCCA GTTTTATATC AGTGCATTAA AGTGCTAAAAC AGTTAGAAA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGTCTCTGC CGGGCTGCA GCACTGCCTA GCAACTTCTT TGGTGGTTGG GTATGAAGT CTCTTCAACG CTTCCTCTCC CTTCCACCGG AACCGAAAAT GTAATCTTAAT ACATATCTAA AATACTATT GCTCTTCAAC GTCCTTAAA AATACTATT TTATTTAATT ACATATCTAAT ACATATCTAAT TATTTAATT ACATATCTAT TTGTTTTGT TAGTTTCTAC GGTGTTCTAA CATGACCAAA AGCACTTTG GGTGTTGTAA CCCCTAAACT TCATGCGTTT TCTTCGGAGT TCTTTCTACC AGGTAGTGTC ACACCGTAC	120 180 240 360 420 480 540 600 660 720 780 840 900 960 1020 1140 1200 1320 1380 1440 1500 1560 1620 1680 1740 1800 1920 1980 2040 2160
50556065707580	GAGCAACCTC CGACCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCAGGGGCTG TGACTCCTTG CATCCTCTG CATCTCTTGAAGAC TCTTGCAGGT ATTCTATGAC GGAAAAACAC GGACATTAGAC TATATTATACT TATATATACA TCATATTGAT CCATATTGAT TCATTGAT TTCATTGAT TTCATTGGT AGCCAAGAAA TTTCATTGT TATATTTTTA TTTCATTGGT TATATTTTCT TATATTTTCT TATATCTTCC GATAACTTT TATATCTTCC GATAACTTT TATATCTTCC GATAACTTC TCATTTGT TATATTTTCC TTATTTTCT TATATTTTCC TTATTTTCC TTATTTTCC TTATATTTCC CTCATTTGT TTTATTTCC CTCATCAGTC CTCATTTCC CTCATCTCC CTCATCTC CTCATCT CTCATC CTCATC CTCATCT CTCATCT CTCATCT CTCATCT CTCATCT CTCATCT CTCATCT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG GCACCTTCG CCTGAGCCAG GCACTTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC ATACTATCAT ACAAAACAAA	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG CCTATCCCGG GCAGCACATT CAATCTTTGT AGAAGATCAG TAGTTGCCAC CAGTCAATGC CAGTCAATGC CAACACCAAG TAACATCAG TAACATTTGT AAACATTAGG CAAACAAACA ATTATCTTTT ATACTTAAAT ATCTTCATTA ATCTTCATTA TCTTCAATT ATAGCACTAG TAGATCAGAC TAGATCAGAC TCCACACA TAGATCAGT TTTAAACATTA TTCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCAATTGAGT TTTAAGCT TTTAAGCT TTTAAGCT TTTTAAGCT TTTTAACTT TCTTCTCTTT TCTTCTCTTT TCTTCTCTTT TCTTCT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC CGACACATC CGACCACGT CGACCACGGG GCAACCACGT GACCACGGT GACACCGGG GCAAGCAACC GGCCACCGTT CAGGTACGAA CAGGTACGAA ACATCGAAA ACATCAAATCA ACCTTAGAAT ACTCAAATGG TCTATTAAA TTTCTTAAA TTTCTTAA TTTGGGGG CATCGTTATT TCACATTTCATA TTTGGAGGCA ACCCGTCATCA ACCTGACA ACCTGACA ACCTGACA ACCTGACA ACCTGACA ACCTGACA ACCTGACA ACCTGAACAA GCTGAACAA GCTGAACAA GCTGAACAA GCTGAACAA GCTGAACAA GCTGAACAA CTTTGAACAT TTTGAACAT CAATTCATA CTTTGAACAA CATACTTAAC CTTTGAACAA CATACTTAAC CTTTGAACAA CATACTTAAC CTTTGAACAA CATACTTAAC CTTTGAACAA CATACTTAAC CTTTGAACAA CACAGTCTAT CTCTCTCTAC CCACACTCTA CTCTCTCTAC CCACACTCTA CTCTCTCT	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGT GGCATCAGA TTTGGTCAGA TTTGGTCAG TTTGAAACA TTTGGTTAAACA TTTGGTAAACA TTTGGTAAACA TTTGGCAC GCGATGAAGT ATAGGTAAT CCTTTGCAC GCCATTTCA AGCCCTTATA TCTTTGACACA TCTTGACCAC TGTTCACAT AATCTTTCTG TCTGACCAT AATCTTTCTC TCTGACCAT AATCTTTCTC AGTGTAATT AATCTTTCTC AGTGTAATT AGTGCTAACA AGTCACTTAA AGTGCTAACT AATCACTTAA AGTGCTAACT AATCACTTAA AGTGCTAACT ACTTACACC AACAAAACCT TTCCACCGA AACTATGCCT TTCCACTGA AACTATGCT TTCCACTGAA ACATATGCT TTCCACTGAA ACATATGCT TTCCACTGAA ACATATGCT TTCCACTGAA ACATCTATTT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGCTTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTCAC GTTCCTCCC CTTCCAGCGG AACCGAAAAT GTAATCTCAA AATACTAT GCTCCTTAAA AATACTAT GCTCCTTAAA AATACTAT TGATTTTAAT TTATTTAAT TTATTTATT TTGTTTTTGG TAGACTAGC TAGACCAAA AGCACCTAC CCACTAAACT TCATGCGTT TCTTCTGGAGT TCATGCGTTT TTCTTCTGGAGT TCATGCGTTC ACGACCGTAC CCAAAACCTAC CCAAAACCTAC CCACAGACCA CCACTGAACCA CCACT	120 180 240 360 420 480 540 600 660 720 840 900 960 1020 1140 1220 1380 1440 1560 1560 1620 1680 1740 1860 1920 1980 1980 1980
505560657075	GAGCAACCTC CGACCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCCCCAGTGG CCAGCGGTGT CATCCTCTG CATCCTCTG CATCCTCTG CATCAGAGA ACTCATATGAC GGACATTGAG GTATGGTATTACT TATATATAGA CTCATTATGT CCATATTGAT TATATTTTTAT TTTCATTGGT TAGCTAAAAAT TTTCATTGGT TAGCTAAAAT TTTCATTGGT TAGCTAAAAT TTTCATTGGT TATATTTTTT TAGCTTTTAT TTTCATTGGT TATATTTTTT TAGCTTTTAT TTTCATTGGT TATATCTTCC GATAATCTGT TATATCTTCC GATAATCTGT TATATCTTCC GATAATTAGT TATATCTTCC GATAATTAGT TATATCTTCC GATAATTAGT TATATCTTCC GATAATTAGT TTTATTTGCT TATATCTTCC AACACTTACT AAACCTACGC	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG CCTGAGCCAG GCACCTTCG CCTGAGCCAG GCACCTTCG CCTGAGCCAG GAGTGTTCC CTGAATCTGA GAGTGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTAC ACACTCTTACC ATACTATCAT ACAAAACAAA TAATCTTATT TGCTTCCCATT TATGTATATA CCATAATCTT CCTTATCTC AAGGATGATT CCATAATCTT CTCATTTACT CAGTTGACC AATACTTTC TCATTTACTC TCATTTATTAC CCTGTTGACC AATATTTATTA TGATTGATT TTGATTGAC TTGATTGAC TAATAGGTG TGACAAATAT ATCTGCCAAA ATCTCGCCAAT TCACTGCCTT CAGTTGCCT TCACTGCCT TCACTCC TCACTC TCACTCC TCACTC TCACT TCACTC TCACT TCACTC TCACT TCACTC TCACTC TCACT TCACTC TCACT TCACT TCACTC TCACT TCACT TCACT TCACTC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG CCTATCCCGA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC TAGTCATTAG TAACATCAG TAACATCAG TAACATCAG TAACATTAG TAACATTATT TACATTATAT TACATTAAAA ATTGGTATAT TTCTTCAATT ATAGCACTAG TAGATCAGAC TTATCAGTTT TTCAATT TTCATTAATT TTCTTCATTA TCTTCAATT TCTTCAATT TTCAATT ACTCTCATT AGACACTGAA TCCTCTCT TTCATTC TTC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC CGACACATC CGACCAGTC CGACCACTC CGACCACTC CGACCACCTC CGACCACCTC CGACCACCTC CGACCACCGT CAGCACCGGG GCAACCACC GCCACCGTT CAGCTACCA ACCATGGTAT CAGCTACCA ACCATTACAAT TCCTCATAT ACATCAAATGG TCTATTATAA ATTCCTTATC ACTTCATATT ACATTACTTTTC CTTCTGGTG CTTCAGACC ACCTGTACC ACCTGTAC ACCAGTCTAT CTCTCTAC CGTGCCTTCC CGTGCCTTCCT CGGCCCTTCCT CGACCACC CGTACCTCC CGGCCCTTCCT CGACCACC CGTACCTCC CGGCCCTTCCT CGACCACTCC CGGCCCTTCCT CGACCACTCC CGGCCCTTCCT CGACCCC CGACCTCCT CGACCCC CGACCCTCC CGACCCTCC CGACCCC CGACCCC CGACCCC CGACCCC CGACCC CGACC CGACCC CCACC CGACCC CCACC CCAC	CCGGGCGCGG CCACACCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT GTGACTTGA GGCATGAAGT ATTGGGGGT ATTGGGGGT ATTGGTCAGA TTTGGTCAGA TTTGGTAAAAT TTTGGTAAAAT AGGGAAGGGT ATAGGAGGTAAT GCTTTTCACAC GCCTTTTCA AACCTTACA GCCCTTTTCA AGCCCTTTT TCTTTGCCAC GCCTTTTCA AACCTTATT AACTTTCTACAC GTTTAAAAT ACGTTACATT AACTTTACAC TCTTACACA AACACTTACA AACACTTACA AACTTACACA TTTTCCCCCA GTTTTATATA AGTTCACTAAA CAGTTAGAAC AACTACACT AACAAAACCT TTCCACTGAAA CAGTTTATTC TCCACTGAAA CAGTTTATTC TCCACTGAAA CAGTTTATTC TCCACTGAAA CAGTTTATTC TCTCCCCCAAAAACCT TTCCACTGAAA CAGTTTATTC TCTCCTCACCA ACAAAACCT TTCCACTGAAA CAGTTCTATTC TCTCTCTCCCCAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTTCTGC CGGGGCTGCA GCACTGCCCT AGGCCATGTT TGGTGGTTGG GTATGAAGT CTGTTCAAGA CTCTCTTCAC GTTCCTCAC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT GCTCCTTAAA AAATACTATT GTATTTAATT ACATATGTAAA AAGACCTAGC TATACTTAT TTGTTTTCTG TAGTTTCTAA AGGACTACT CATGCGTTT TTCTGGGGT TCTTCATCCAC GGTGTTTTAC CCCTAAACT CTCATGCGTT TTCTTGTGT TTCTTGGGGT TCTTTCTACC AGGTAGTGT ATTTTCACC AGGTAGTGT ATGTAGTTT CATGCGTTT TTCTTGAGGT CCTTTCACC AGGTAGTGT ATGTAGTGT ATGTAGTGT CCACACGTAC CAAAACCTAC CCACTGAACA GTCTATTTCC	120 180 240 360 420 480 540 600 660 720 840 900 960 1020 1140 1220 1380 1440 1500 1560 1620 1680 1740 1800 1860 1920 2040 21100 2220 2280 2340
50556065707580	GAGCAACCTC CGACCAGAG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG CCAGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCAGAGA ACTCATCATGAC TCTTGCAGGT ATTCTATGAC GGACATTGAG GTATGGTATT AAACATGCT TTGTATACT TTGATTACT TATATATAGA CTCATATTGAT CAGTCAAATTA TTTCATTGGT AGCCAAGAAG GTAATATTAC TTATTTTTTA ACCTTTTTG ACCAACTTTA ACCTTTTTG TATATATTGCT TATATTACT TATATTACT TATATTACT TATATTACT TATATTACT TATATTTTC GATAATCTCC GATAATCTG TCTTTTTTC TATATTACT ACCTACTGA ACCATACCT AAACCTACCT AAACCTACCT AAACCTACCT	AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TCATTCTCCG AGGATTTACT TGGATGTCTC CTGAATCTGA GAGTGATAG GCTAGTTC CCTGAGTATT CCTATGACCC GTGTGTTTC ACCTCTTACC GTGTGACACA ATACTATATT TGATTCATT TAGTTATATA TGATACTATT TCATTTACTC AAGGATGATT TCATTTACTC AAGAACAA CCCATTTTCCCATT TCATTTACTC AAGAATGTTT TCATTTACTC AATATTATT TCATTTACTC TCATTTACTC TCATTTACTC TCATTTACTC TCATTTACTC TTAGTTGACC AATATTTTT TCATTTACTC TCATTTACTC TCATTTACTC TCATTTACTC TTATTATATACC TCATTACTC TTAATAAGGTG TGACAAATAT ACCTGCCATT TCACTGCCTG TCACTGCCTT TCACTGCCTT TCACTGCCTT CCTGTTCTCC TCATTGCCTT TCACTGCCTT CCTGTTCTCC TCATTGCCTT ACATACCTTC	ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCAGCACATT CAATCTTGT AGAAGATCAG CAGCACATT CAATCTTGT AGAAGATCAG AGGCAAAG CAGCAAAG CAGCAAAG AACAAAAAA ATTACTTCTT GAGTAATCATT ATACTTTCATT ATACTTAAAA ATTGGTAATT TTCTTCAATT ATAGCACACA CAAATCAGAC CAAATCAGAC CAAATCAGAC TTACTTCATT TTCTTCAATT ATAGCACAC CAAATCAGAC CAAATCAGAC CAAATCAGAC TTCCACACA CCAATTGAGT TTTTAAGCTAA TTGATCTTTT TCTCTCTGTA TTGAGATATT TCTCTCTGTA TGGCCTTC CAGTGCCTTC ATGTGCCTTC ATGTGCCTTC ATGTGCCTTC ATGTGGCCTTC ATGTGGCCTTC ATGTGGCCTTC ATGTGTGTTTT	CAGCGCCGCC CGAGCAGGGC CTTGCCCACCT CGAGCAGGTC ATGGATCGGC GGACACACTC GAGCACCGTT GATGCATCC GAGCACCGTT GATGCTATC GAGCACCGTT CAGGTACGAA GGAGAGTACCAA GGAGAGTACCAA ACCTTAGAAT ACAAAAACCCAT TCCTCAAAAT TCCTCAAAAT TTCTTTTTC GCTTTGGGTG CATCGTTAT ACATTCATA TTTGGAGGCA ACCTTACATCAT TTTGGAGGCA ACCTTATCATA TTTGGAGGCA ACCTTACATCAT TTTTCTTTTC	CCGGGCGCGG CCACACCT TCCCACACC TCCCACACC GCCATCGCCC CCAGATCCACG GCCATCGCCC CAGATCCACG GCCATCGCCC CAGATCCACG GCCATCGCC CAGATCCACG GCCATCGCT ATTGGCGGT TTTGGTCAG TTTTGGTCAG TTTTGGTAAAA TTTGGTAAAA TTTGGTAAAA TTTGGTAAAA TTTGGTAAAA TTTGGTAAAA TTTGGTAAAA TTTGGTATT GCCTACACT CCTTTTCAC CCCTTTTCA AAGCCCTTAT GCCCTACATT TCTACACCA TTTTACACC TTTCACAC AGTCATTAA AGTGCTAAA ACTTTAGAA ACTTTAGAAA ACTTTAGAAA ACTTTAGAAA CAGTCAATT TCCACTGAA CAGTCTATT TCCACTGAA CAGTCTATTC TCTCACCC TTTCCACTGAA CAGTCTATTC TCTCTACCC TTTTTAACAAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGTCTCTGC CGCGCTTCTGC CGGGGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTCAC GTTCCTCCC CTTCCAGCGG AACCGAAAAT GTAATCTCAA AATACTAT GCTCCTTAAA AATACTAT GCTCCTTAAA AATACTAT TGATTTTAAT TTATTTAAT TTATTTATT TTGTTTTTGG TAGACTAGC TAGACCAAA AGCACCTAC CCACTAAACT TCATGCGTT TCTTCTGGAGT TCATGCGTTT TTCTTCTGGAGT TCATGCGTTC ACGACCGTAC CCAAAACCTAC CCAAAACCTAC CCACAGACCA CCACTGAACCA CCACT	120 180 240 360 420 480 600 -660 720 780 840 900 1020 1140 1260 1320 1320 1340 1560 1680 1740 1860 1740 1890 2040 2100 2220 2280 2340 2400

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2520
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       AGCAAGGCAT TTGGCTGCTG TAAGCTTATT GCTTCATCTG TAAGCGGTGG TTTGTAATTC
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       CTGATCTTCC CACCTCACAG TGATGTTGTG GGGATCCAGT GAGATAGAAT ACATGTAAGT
                                                                          2640
       GTGGTTTTGT AATTTGAAAA GTGCTATACT AAGGGAAAGA ATTGAGGAAT TAACTGCATA
                                                                          2700
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                                                                          3000
10
                                                                          3060
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       CTACACAAGG AAAGTCAGCC ACCGTGTCTT ATGAGGAATT GGACCTAATA AATTTTAGTG
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       TGCCTTCCAA ACCTGAGAAT ATATGCTTTT GGAAGTTAAA ATTTAAATGG CTTTTGCCAC
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30
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       Coding sequence: 897-1400
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       ATAGAGGGAA TGAGTATTAA TTGGAGAAGC TTAAAGTATT GCCACTTTAG CACTGAAGAT
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55
       TGGGATGAGA GGAGGTGAAA CCTCACTAGA AAAAGGGACA ATGTTAGTGT GGCCCTTCCT
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                                                                           420
      GTTGAGTGTA TACTGTCTGT CAAAGACTTC CAGCATTTCC AGGTCCTAGA GAGGAACAAG
ACTGGTAACC TGCCTATCTG TATTTTTAAG AACCCAGGAG GAAAGCTTTA TAATAGAACA
                                                                           480
                                                                           540
       TTATTTCTGT GTTTATGTAT AAGGGGTTTT TTGTTTTTTT AAAGACAGGA TCTCACTCCA
                                                                           600
60
      TTGTCCAGGC CAAGTGCAAT GGCACGAACC TCATAGCTCC TGGACTTAAG TGATCTGCCT
      TAGTCTTGCT TTGTTGCCAG GCTAGTCTCA AACTCCTGGC TTCAAGTGAT CCTCCTGCCT
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       CAGCCTCCCA GAGTGCTAGG ATTACAGCAC TTGGATTCAG CTTCTTCATT TCCAACATGG
                                                                           900
65
       AAGAAACTTA CACCGACTCC CTGGACCCTG AGAAGCTATT GCAATGCCCC TATGACAAAA
                                                                           960
       ACCATCAAAT CAGGGCTTGC AGGTTTCCTT ATCATCTTAT CAAGTGCAGA AAGAATCATC
                                                                          1020
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                                                                          1080
       CTGAAATTAG TCATCATATC TCAAGCTGTG ATGACAGAAG TTGTATTGAG CAAGATGTTG
                                                                          1140
       TCAACCAAAC CAGGAGCCTT AGACAAGAGA CTCTGGCTGA GAGCACTTGG CAGTGCCCTC
                                                                          1200
70
       CTTGCGATGA AGACTGGGAT AAAGATTTGT GGGAGCAGAC CAGCACCCCA TTTGTCTGGG
                                                                          1260
       GCACAACTCA CTACTCTGAC AACAACAGCC CTGCGAGCAA CATAGTTACA GAACATAAGA
                                                                          1320
       ATAACCTGGC TTCAGGCATG CGAGTTCCCA AATCTCTGCC GTATGTTCTG CCATGGAAAA
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       ACAATGGAAA TGCACAGTAA CTGAATACCT ATCTCATCAA ATGCCAGACC CTAGAAGACT
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       GITGCTTCTT CTTCTACCAG TGGGTTCTCA TTTTCCTCCT AATCTAATTA TAGAATGGTA
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       AACTCCCTGT GACTTTCCAA ACTGACAAGC ACACTTTTTT CCTCCCCCCT TGAATCCTCA
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       RAEISHHISS CDDRSCIEQD VVNQTRSLRQ ETLAESTWQC PPCDEDWDKD LWEQTSTPFV
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WO 02/086443
Seg ID NO: 248 DNA sequence
Nucleic Acid Accession #: NM_003392
Coding sequence: 758..1855

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	TTDAGGGAAAT	CCGGGCTGCT	CTTCCCCATC	TGGAAGTGGC	TTTCCCCACA	TCGGCTCGTA	60
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	CGAGGCCGGC	CGCAGGACGG	TGTACAACCT	GGCTGATGTG	GCCTGCAAGT	GCCATGGGGI	1500
	GTCCGGCTCA	TGTAGCCTGA	AGACATGCTG	GCTGCAGCTG	GCAGACTTCC	GCAAGGTGGG	
	TGATGCCCTG	AAGGAGAAGT	ACGACAGCGC	GGCGGCCATG	CGGCTCAACA	GCCGGGGCAA	1560
	GTTGGTACAG	GTCAACAGCC	GCTTCAACTC	GCCCACCACA	CAAGACCTGG	TCTACATCGA	1620
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	CCCACCCACC	ACTCAGCCCC	GCTCCCAGGA	CCCGCTTATT	TATAGAAAGT	ACAGTGATTC	1920
40	TO COMPANY OF	ተሞተሞተልርልልል	ተልተሞሞሞሞ AT	TTTTCCCCAA	GAATTGCAAC	CGGAACCATT	1980
	ጥጥጥጥጥጥርጣር	ጥጥልርርልጥርጥል	AGAACTCTGT	GGTTTATTAT	TAATATTATA	ATTATTATTT	2040
	CCCAATAATC	GGGGTGGGAA	CCACGAAAAA	TATTTATTTT	GTGGATCTTT	GAAAAGGTAA	2100
	TACAACACTT	CTTTTGGATA	GTATAGAATG	AAGGGGGAAA	TAACACATAC	CCTAACTTAG	2160
	CTCTCTCCCA	CATGGTACAC	ATCCAGAAGG	TAAAGAAATA	CATTTTCTTT	TTCTCAAATA	2220
45	サウクウカ かつみ すみ	TOGGATGGGT	AGGTTCCAGT	TGAAAGAGGG	TGGTAGAAAT	CTATTCACAA	2280
	ምምር እርያርጥምርጥ	ΑΤΓΙΑΓΓΑΑΙΑ	TGAGTTGTAA	ATTCTCTGGT	GCAAGATAAA	AGGTCTTGGG	2340
	מממממממת		AAACCTCCCT	TCCCCAGCAG	GGCTGCTAGC	TTGCTTTCTG	2400
	CATTTTCAAA	ATCATAATTT	ACAATGGAAG	GACAAGAATG	TCATATTCTC	AAGGAAAAA	2460
	CCTATATCAC	ልጥርጥርጥርግልጥጥ	CTCCTCAAAT	ATTCCATTTG	CAGACAGACC	GTCATATTCT	2520
50	N N TO N C C T C N T	CALATTTCCC	CAGCAGGGAG	GAAAGTCCCC	AGAAATTAAA	AAATTTAAAA	2580
	CONTRACTOR A TRUTTER	ADCIDECTA	TTTGAAGCTG	TTATAAGAAT	TGGGATTCCA	GATTTGTAAA	2640
	አ አ ር አ ር ር ር ር ር ር ል	ልጥናልጥጥርፕር ር	ACACTAGATT	TTTTGTTTGG	GGAGGTTGGC	TIGAACATAA	2700
	ΑΤΓΙΑ ΑΑΤΑΤΟ	CTGTATTTTC	TTAGGGATAC	TTGGTTAGTA	AATTATAATA	GTAGAAATAA	2760
	TACATCAATC	CCATTCACAG	GTTTCTCAGC	CCAAGCAACA	AGGTAATTGC	GTGCCATTCA	2820
55	CCACTCCACC	AGAGCAGACA	ACCTATTICA	GGAAAAACAG	TGAAATCCAC	CTTCCTCTTC	2880
55	አሮአሮተርአርርር	CTCTCTCATT	CCTCCGTGTT	GTGATGTGAT	GCTGGCCACG	TTTCCAAACG	2940
	CCACCTCCAC	マロススマア アプログラ	TTGGTTGTAG	GACAGGAAAT	GAAACATTAG	GAGCTCTGCT	3000
	TOCABARCAG	ידידים בידים בידידי	AGGGATTTT	GTTTCCTAAA	ACTTTTATTI	TGAGGAGCAG	3060
	ጥ አ ውጥጥጥጥ ውጥ አ	ጥር ጥጥጥ ላ ልጥር	ACAGAACTTG	GCTAATGGAA	TTCACAGAGG	TGTTGCAGCG	3120
60	TATION CTCTT	ከጥር ከጥር ርጥር ጥ	CTTTACATTA	TCCACTCATG	CTTCTCCTAT	TGTACTGCAG	3180
00	CONTRACCORD N	ል እ እ ርጥርያጥጥር ር	СУСТСТУСТТ	GAACAGTTGC	ATTTATAAGG	GGGGAAATGT	3240
•	ርርምምያ እንተርር	ሚያርር የሚያው የሚያ	CTCAAAGTCT	' TTTGTACATA	. ACATATATAT	ATATATACAT	3300
	TARATATA	Α ΤΑ Α ΤΑ ΤΑ ΤΑ Α	ATATATCTCA	TTGCAGCCAG	TGATTTAGAT	TTACAGCTTA	3360
	WINIVIUM	ስተርቀርቀርቀር የደረፈተር ነው	CTAGAGCATT	GTTGTCCTTC	ACTGCAGTCC	AGTTGGGATT	3420
65	ATTCCANANC	TTTTTTTCACT	CTTGAGCTTG	GGCTGTGGCC	CCGCTGTGAT	CATACCCTGA	3480
05	CCACCACCAA	GCAACCTCGT	TTCTGAGGAA	GAAGCTTGAG	TTCTGACTCA	CTGAAATGCG	3540
	TOTAL CONTROL	* እስርአጥአጥርጥጥ	լունդեր և դորդորը <u>։</u>	CTGCCTCACC	CCTTTGTCTC	CAACCICCAI	3600
	TGI IGGGI IG	THE TREE PROPERTY.	GGGCATTACT	TGTTCGTTAT	AGACATGGAC	GTTAAGAGAT	3660
	TICIGIICAC	, IIIGIGGAGA	ACCANTETT	CTCTTTTCTT	AGTTCATTCT	GCAGAATGGA	3720
70	ATTCAMARCI	CHUMMUCAIC	CACACTACTA	מבטדדם בידים	TCCCTAAGGA	ATATTCAGCC	3780
70	AACCCAIGCC	, INIINIMANA		V dalalalalalalata V dalalalalalalata	TANGGACACC	TCTTTCCAAA	3840
	CACTACATAC	, WINGCITTI	ሚአጥርጥርክርልር	ጥተልተርተዋርሞባ	TTAAAAGTTT	GGAAAGATAC	3900
	CAGGCCATCA	AMINIGITOI	TAICICAGAG	CCCCTTTCAT	ATCACCTCAC	CCAACTGTGG	3960
	ACATCTTTTC	. AIACCCCCCC	. IIMGGAGGII	TCACCCAACT	GTGGCTCTTT	AATTTATTGC	4020
75	CTCTTAATTI	. ALIGCATAAT	- ANTHIOCHOL	TOTOCOMO	GCAAAAGATO	TTGAAAGCAA	4080
13	ATAATGATAT	TCACATCCCC	, ICAGIIGCA	ուսարությունը և հետևարարական արգագրարարան անագրարարացում է արարագրարան արարարարան արարարան արարարան արարարան ա	ATTATACAA	AACCATGAAG	4140
	AAAGCACTAF	TAGTTTAAA	MIGICACITI	· Controporting	GACTCATGT	TATGAAGAGA	4200
	TACTTTTTT	ATTIGCTAAA	TUAGATIGIT		ATCTABATT	TTCTACATGT	4260
	GTTGAGTTT	ACAATCCTAG	CITTTAAAAU	հ մահիմահանանան Հ ԿԱԿԿՐԴԿՐՐՐ	, առագողություն Մարդություններ	ACATATTTCT	4320
80	CATTCAGATA	TTATGTATAT	CTTCTAGCC	LIMITOTOL	ACDTOCADA	GCTTATTCCA	4380
οU	GTCTTGCGTC	ALITGIATAT	TTCACTGGT	· INTERNATION	. COLOGRAM		
	AATGGAAGAT	r agaatataa	ATAAAACGT	ACTIGIAAA	· constituted and a second a second and a second a second and a second a second and		
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85	Profesu W	ccession #:	W&_003383				
0)	•		23	31	41	51	
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MAGSAMSSKF FLVALAIFFS FAQVVIEANS WWSLGMNNPV QMSEVYIIGA QPLCSQLAGL
                                                                               60
       SQGQKKLCHL YQDHMQYIGE GAKTGIKECQ YQFRHRRWNC STVDNTSVFG RVMQIGSRET
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       LADFRKVGDA LKEKYDSAAA MRLNSRGKLV QVNSRFNSPT TQDLVYIDPS PDYCVRNEST
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       TCGGCCAGAT GTGGTGAGGG CTAGGAAAAG AGTTTGTTGG GAACCCTGGG TTATCGGCCT
       CGTCATCTTC ATATCCCTGA TTGTCCTGGC AGTGTGCATT GGACTCACTG TTCATTATGT
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       TCAGGTTATC AAGTTCAGTC AACAGAAGCA TGGAGTGTTG GCTCATATGC TGTTGATTTG TAGATTTCAC TCTACTGAGG ATCCTGAAAC TGTAGATAAA ATTGTTCAAC TTGTTTTACA
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       GCCTGGTGTT TATACTAGAG TTACGGCCTT GCGGGACTGG ATTACTTCAA AAACTGGTAT
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       CCATTTTTAG AGATACAGAA TTGGAGAAGA CTTGCAAAAC AGCTAGATTT GACTGATCTC
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       DKLYAEFGRE ASNNFTEMSQ RLESMVKNAF YKSPLREEFV KSQVIKFSQQ KHGVLAHMLL
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       RSKTLGQSLR IVGGTEVEEG EWPWQASLQW DGSHRCGATL INATWLVSAA HCFTTYKNPA
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       RWTASFGVTI KPSKMKRGLR RIIVHEKYKH PSHDYDISLA ELSSPVPYTN AVHRVCLPDA
                                                                              300
       SYEFQPGDVM FVTGFGALKN DGYSQNHLRQ AQVTLIDATT CNEPQAYNDA ITPRMLCAGS
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       GAGGGTCCTT CTCTTCGTGG CCTCGGACGT GGATGCTCTG TGTGCGTGCA AGATCCTTCA
                                                                              180
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                                                                              300
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                                                                              360
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                                                                              480
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       GGAGGCCCGG AGAAGAGACA TCCTCTTTGA CTACGAGCAG TATGAATATC ATGGGACATC
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10	Seq ID NO: 253 Protein sequence: Protein Accession #: NP_003495.1						
15	AFLEHKEQFH QDDDLEVPAY RRDILFDYEO	11 YEVVQSQRVL YFILINCGAN EDIFRDEEED YEYHGTSSAM RHNHRNEDEE	LFVASDVDAL VDLLDILQPD EEHSGNDSDG VMFELAWMLS	EDTIFFVCDT SEPSEKRTRL KDLNDMLWWA	HRPVNVVNVY EEEIVEQTMR IVGLTDQWVQ	NDTQIKLLIK RRQRREWEAR DKITQMKYVT	60 120 180 240 300
20	KLWSVHGQKR FSIHFGFKHK KQLRATQQTI TKNRRCKLLP	LQEFLADMGL FLASDVVFAT ASCLCTNLVI LVMAAPLSME AEDRSKFLDA	PLKQVKQKFQ MSLMESPEKD SQGPFLYCSL HGTVTVVGIP	AMDISLKENL GSGTDHFIQA MEGTPDVMLF	REMIEESANK LDSLSRSNLD SRPASLSLLS	FGMKDMRVQT KLYHGLELAK KHLLKSFVCS	360 420 480 540
25	Seq ID NO: 254 DNA sequence Nucleic Acid Accession #: NM_022337 Coding sequence: 48683						
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35	TCATCAAGCG ACTTCGCGCT ATATCGCAGG GTGCATTTAT AAAATGATTT	CTACGTGCAC CAAGGTGCTC TCAAGAAAGA TGTCTTCGAT GGACTCCAAG	CAGAACTTCT CACTGGGACC TTTGGAAACA GTCACCAGGC TTAAGTCTCC	CCTCGCACTA CGGAGACTGT TGACGAGGGT CAGCCACATT CTAATGGCAA	CCGGGCCACA GGTGCGCCTG CTATTACCGA TGAAGCAGTG ACCGGTTTCA	ATCGGCGTGG CAGCTCTGGG GAAGCTATGG GCAAAGTGGA GTGGTTTTGT	180 240 300 360 420
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		267 Proteir ession #: >					÷
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WO 02/086443

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Coding sequence: 247-816

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0 0	1	11 	21 	31 	41 	51]	
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WO 02/086443

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WO 02/086443

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.0		COURT ACCENCY	CCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCC	CACCCCACCT	GACACGGAGG	COCIOCICIO	240
		CCCTCCTACA	CCCCCACTC	CCCCAAGCIG	AGLGCIGCCG	AGGCCCAGAA	300 360
	443 AMAGA 34	CACAACCCCC	CCV VCCTCCC	CACTGTGAAG	AGCAAGGAGG	AGGCCCAGCA	420
.5		N BORGOCCECC	AGCTCCTGAG AGCGAGAGAA	CCCCAAGTGC	CTGGACCCTA	GICIGCOGCI	480
.3	a aaaaamma	NO COMPOCIONO CO	CCCCCCCCCA	GGACACGCCT	TACTCTAACT	GGCACAAGGA	540
	00m00000xx0	TO THE PROPERTY OF THE PROPERT	CCV VCCCCCCC	TGTGTCTCTG	CIGCIGGACC	IGICCCWGCC	600
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١٥	CGGAAGTAAC	ATTGAGGGCT	TCGTGTGCAA AGGTGACCTA	CACCACCCCC	TTCCAGACCA	CCAGTTCCTC	780
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		A CAMPA MALA LALL	ጥርርጥርጥርር ልል	GGAGAAGGCC	CCCGATGTGT	TUCHCIGGG	900
	~~~~~~~~		<b>サクトのクククククス</b>	GTATGGCTGC	AACTTCAACA	AIGGGGGCIG	960 1020
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	A A . CCC . C	CACAACAACC	ACAAGAAGCC	CCAGAATGCG	GCAGACAGTT	ACTUCTOGGI	2040 2100
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45	G3 3 3 G4 3 DOC	ጥ አ አርጥርጥርርጥ	CCTTAAAGGC	CCCTTGGAAC	ATGCAGGTAT	TTTCTACGGG	2280
		かいしゅう かんかん	CANCOTOTOT	CTTCCCCTCC	CACGGTGGGG	ATTICGICAC	2340
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50 .	COMP. CCP. TCP.	ስ ከ ስ ርጥል ል ል ጥር	ከልጥጥልልጥጥል ፕ	TCAATTAGGI	AAGAAGATCI	GGIIIIIIGG	2580
JU ,	TO	ことのことを	CTCCAAACAT	TTCTTTACAI	TTGCATTCC	CCATTTCGCC	2640
	3003030EC	መምረም የሚያ ነው	ጥር እጥ እርጉርጥጥ	GACATCCTCC	AGAATGGCCA	GAAGIGCAAI	2700 2760
	TAACCTCTTA	GGTGGCAAGG	AGGCAGGAAG	TGCCTCTTIA	A ACTTOTTACE	TTCTAATAGC AAGTGCATTA	2820
55	as concerno	サース カーサース へん	ተልልተርቸልርርር	GGCTAGGGCG	AGAGAGGCCA	GGGATTTGTT	2880
33	AR AR ARER OF	የመጣ አለጥጥ አለጥጥ	<u> </u>	TACTGAGGTI	ACCACACAC	TURCIACUM	2940
	mamas mass se	• እርጥእእር <u>እ</u> እርር	האבעמבעמים בייבור ב	AAGGACAACC	TGTCTTTGAG	CCAGGGCAGG	3000
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60	## ## ## ## ## ## ## ## ## ## ## ## ##	でみつずむであずなど	• አርር አጥርር ፕርጀ	GATTTTAAAI	CCIGAAGIG.	GGGTGGCGCW	3240
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	OTO CTTT A A A	ጥጥ/?!!!!!!!! የተመሰው የ	CCACCGAAGG	: AAGAGGGAAA	\ GAGATGACT/	A WCIMMMICH	3360 3420
CF	TTTTTACAGO	AAAAACTGCT	CAAAGCCATT	TAAATTATA	r ccrcarrii	AAAGTTACAT TCTCTCTCTC	
65	mamamana a	* ***********	፣ አሮአሮኔሮኔሮኔር	י אראראראני	: AGAGACACG	CACCATICIO	3240
	aamaaaaaa	3 መሮሮኤክሮ እሮ እባ	• • • • • • • • • • • • • • • • • • • 	" CACCGATGG"	r CAGAGTCAC	AGAAGIIACC	3000
	man ama mana	~ ~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	י כאיניתכיתכית	r GTGGGCTTT.	I TACCACCAC	I GIGCHGGHGW	3000
	10101010101	ግ ሮ አአአምርምርም	፣ ጥርርር የነርር አ ር	R GCCCCAAAG	C CTCAGAGAA	A GGGTGTTTCT	3120
70	GGTTTTGCC	r TAGCAATGC	A TCGGTCTCTC	AGGTGACAC	r CIGGAGIGG A CATCCTATG	TGAAGGGCCA TTCAGATTGT	3840
	COMPAN A STACE		A GGCAGGGGAI	A CTCARAGGA	A AGATGGAGG	T TTTGTGCGGC	3300
	maan maaaaa	~ ATTTCCAAC	r ጥርጥጥጥጥጥልል	A GTCATCTCA	T GGTCTCCAG	T TTTCAGTIGG	3900
	* * ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	_ manananana	T NACCCACACAC	A AACCCTGTG	T CCATTIGGC	W WWWCTICCII	4020
75	000010010	A CALCADA CCACA	<u>አ ጥርጥርንፕር</u> አልርያ	~ TGGGCAGTC	T GTGGTGTGG	A GAGCAGCCAI	4000
	CTGTCTGGC	C ATTCAGAGG	A TTCTAAAGA	C ATGGCTGGA	T GOGGLIGGIG	A CCAACATCAG C CTTGCCCTTA	4200
	max mmmcccc	<u>ጥ ጥሮ</u> አለርርአርአ	רויייייייייייייייייייייייייייייייייייי	C TTGGCTTCC	C ACAGCCCCA	W COCHOICIGI	4200
	400 mas mma	~ TOOCONTOON	* ~~~~~~~~~~~	C TATTTCAC	A GIGIICIGA	1 IGCICICACA	4320
80	4444	こ かいしいかしかにか	マ してしてにひひてに	C AGCCCTGTT	C TCAACAACA	G GGWGGICWIG	4300
	CARCOCCEC	T CTCC38CCC	n സമാനാവരെ മ	A AATGGGTGA	T AAAGAATCU	A GIICCICAAA	4440
		a conceence	ש אירירישור באירירי	T CCTGGGTGG	T GCTTTCTC1	T GCACACCACT C ATCTGATGGA	4500
		O CTCSSSCSC	տ ա√ւ∿սեսերը դրդր	ሮ ጥርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርርር	T TTTTGCTG1	T ACTITIONAGE	4020
85	*******	— TTCTTCCC	а сатсттесс	G CTACAGTAG	T GGGTAAAC	A ATGUCUACUG	4000
00	CCCS SCSCC	יה המתחתמה מי	v account cands	T CCTGAGGGG	C CCCAGCTIV	C TUBUSCUTUS	4/40
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21

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	J. J. S. STEPHOOD						

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Seq ID NO: 406 DNA sequence Nucleic Acid Accession #: Eos sequence

WO 02/086443 Coding sequence: 1..927

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			KIFQVVPIPV		·		
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. •	ATCABACTTC	GAGTGCTGTG	CONCAMMINATION 1	ተተርተተር ልርርተ	TCACTGACGG	CCACGGTGGC	60
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10	TTAAATTAT	ATATGTTAAT	AAAAAGGGGG	TCCACAAAAC	AAACAGCCCC	CCACTCTGGT	2340
	TOTCAGGGAG	ስጥስጥጥርርስ TT	AAATGGCCTT	GGACAACAAC	CCCTCTCCCT	GGCCACAGAC	2400
	1G1CAGGGAG	WINTIGGVII	WWW.IGGCCIT	AACACTGGAA	TGAGTCTGAA	CCCAGGTGCT	2460
	ATTCTTCAGA	TIACAAGATA	TICCAGGGGA	MACACIGGAA	TOUGHCHOM	TATTTTTCCC	2520
	AAACAGAAGG	ACCATTGAGA	AATGITGIGA	TCCTGACAGG	TCAAGCAATT	TATTTTCGG	
1.0	CTTCATTTTT	AAATGTAAAA	TTAGAAAGCT	GCCATTTAAA	ATGGCCCGTC	TGTTTCAATT	2580
15	GCTCTTCTCA	GTGTCAGCCT	GTTAACTCAA	TGTGTTAGTC	TGTTTTCATG	CTGCTGATAA	2640
	AAACATACCT	GAGACTGGCA	AGAAAAAGAG	GTTTAATTGG	GCTTAGAGTT	CCACGTGATT	2700
	GGGGAGGCCT	CAGAATCACA	GTAGGAGGCA	AAAGTTATTC	TTACATGGTG	GCTGCAAGAG	2760
	ANGATGAGGA	AGAAGCAAAA	GAAGAAACCC	CTGATAAACC	CATCGGATCT	CCTGAGGCTT	2820
	AMEN'S CHINECOL	ATCACAATAC	CACAACAAAG	ACCGGCCCCC	ልተርልተተርልልተ	TACCTCTACC	2880
20	ATTANCIATE	AIGAGAAIAG	CACAAGAAAG	COMPONENCE	VIOUT TOUR	A CA DOTTCCCT	2940
20	TGGGTCCCTC	CAATAACATG	TGGAAATTCT	GGTAGATACA	ATTCAAGTTG	AGATTIGGGI	
	GGGAACACAG	CCAAACCATA	TCACTCAGCA	AGGCAGAȚAA	CITTCTCACT	GAGCCTATGC	3000
	AACAGAAAAC	CATCTGGGAT	GGTTGTAAGG	GGCACAGĠAA	GTGACTGGTA	GGATCACTGC	3060
	CAAAGCTGAG	CACTCAGGAG	AAGGCAATAG	AATCCTATTC	TCCATAGTAT	GCTATAAGAT	3120
	ACTGAAGTAC	ACTTCTTCAC	TATCTCTTTG	GACTTAGAAT	TAGCACTACA	TTCCTTGTTA	3180
25	TACAGAAAAA	TTACTAACCA	AATTCATAGG	ATGACAAAAA	CTTTCAGAAC	TGAAAAACAG	3240
23	IACAGAAAAA	TIACIANGGA	ANTICATAGO	CGAAGTATGC	CTTTTTTTTT	ATCCAAAATC	3300
	GAAAIGIAAG	CITITIAGIT	CITIOGIAIT	COARGIAIGC	CIMPONGACA	WIGGING III	3360
	CAAGAAAAGA	ATGGTGGGGT	TTTTGTTTGT	TTGGTTTTGT	TITIGITITA	CAGCIGGAGI	
	AGAATACAAA	GGGATGGAGT	TGAAACAAAT	GAGAGGAAAT	TGGAATTCTA	AACTTATTCT	3420
	CATTGGCATT	AGAAAGGCAC	CTACATGTAT	TTCACATGAG	CCGGTGACTG	CTGACTTGCA	3480
30	TTCTTATTTT	TTCCCTATAG	ATTAAAAAGG	AGGTACAATG	GTAGAACTGT	AATCCTGTCC	3540
	ተተተርተርልጥልል	ATTTTCATAT	TCATAAAGGT	GAGTGTTAGC	CCGCTTGTGA	AATCTGAAGT	3600
	TO A COURT A COURT	CANATACTAN	CCACAGAGGG	AAAGGCAGCA	AGAGGAGAGG	CATABATTTA	3660
	IGAGIAACII	CAMMINGIAM	CCACAGAGGG	CAGCCTCTCT	CCCCACCTCT	COUNCOTON	3720
	GGATCTCACC	CITCATTCCA	CAGACACACA	CAGCCICICI	GCCCACCICI	GCIICCICIA	3780
25	GGAACACAGG	TAAGAGCTTC	AAGCCTCTCC	AGCTTAATAA	CATGAATTAT	TTTTGAGAAT	
35	AATAATGATA	CTGTGTTCTA	TATCATGCAT	CTCCTGCATT	CTGTCTGATT	ATATTTTACT	3840
	TATTCTGCCA	GAGCAAAATT	AAAATACCTA	TTTCATCTGA	TTTGTCCTTT	ATCTAAATTG	3900
	CTTAGTTCCA	AGTAAACCAA	GGCACTTTTA	GGAACACAGA	GGGAGAGTGC	CTTGCAGCCA	3960
				TCTTAACAGC			4020
	ACCROTOTIO	THE CONTROL	TTCTA A A CCC	ATCCTACCTA	CCTCTACTCT	AACCAGCAAT	4080
40	AGGICICCIG	TIAGCATICA	TIGIAMAGCC	TTTACAATAG	CCICIACIOI	CONT CONTROL	4140
40	GAAAGAAAGA	TAAAGAGGGT	CGATTACTTA	TITACAATAG	ICITIAAAA	CGIAGIIIIG	
				TAATATATGC			4200
				CAAAATCTCA			4260
	TTTAAGAAAA	TTGTACTACA	AAATACCATT	CCATTTATTA	AAGTCATTCT	GACAGGAATC	4320
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45	CAACACCAAG	TTCATGTTCG	ATCTGTTCCA	ACAGTTCAGA	AAATCAAAAG	AGAACAACAT	4440
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				TATCAGCATC			4560
				GATGGTAATA			4620
50				ACTGTGAGAC			4680
50				TCTCTTCCAG			4740
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	СУССТТТУСУ	TCCCTGAACA	GGTCATAGTT	TAAACCTGGA	ACTTCACAAA	AACTAAGAAA	4860
	ACCCCACTOR	TACCCAAAAT	CTTCCACACA	AAGATTGAGA	CATACAGAGT	CCCTTCCCAT	4920
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	GCCTTGCCCA	TAAAAGTCAT	AATAAATGTT	ATTATTATTA	TAAAGTAGCT	ATAATTATAC	5220.
							5280
60	TWAT COLORS	GWGGY Y CWYW	**************************************	ATCATTCOTA	TOTAL	TGGTTTAGAG	5340
00 ,	ATAAGCACAA	GICCAAGIAI	ATTTTGGAAA	AIGAIIGCIA	TOOUTHINITY I	ACACCOTTAC	
	CCTTAATAGT	GCAAAATGCT	TTGCTGGAAG	GTAGAAAGTT	CTAGATITAA	ACAGGCTTAG	5400
	GTTCAAAACT	TGGCACTTCT	AATTTATGTC	TCTATAAACA	GGGTTTTTTT	CCCCATTCTC	5460
	TGAGCTTTCT	TGTGTTCATC	TGAATTGAAC	TAAAGACTTA	GAGTTACCCA	TGTAAAGTCC	5520
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65	PECCACAGAT	CACTCAATGT	GTCCTACAAG	ATAATAGCAC	CAACAGGTAT	AACAGGGCTT	5640
00	COTCCCATA	TOTATION A A	ATATCCAACC	TTCAACATAC	TOTATOTT	GATGACTGTT	5700
	CCIGGCAIAA	TCINIII INNA	ATATCCANCC	ACCORCACA COT	TTANCTCCCA	AGCTAAACCT	
	AGAAGTGAAA	TATGGTCCTT	GCCCATAAGG	AGCIGAGAGI	TIMACIGOUA	VOCIVILICAT	5700
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=-	AAGAGAAAAG	ATTGCAGTAC	GTTAGAGCAA	GAAGAATTTT	CTGGAAGAAG		5880
70	GGTGGATTTT	GAAGGGTATT	TGAGGTGAAA	TACACCAATT	ATCAGGGAAT		5940
	GTCCTCAATG	AGACTACCAG	CATTTAGGGA	CTGATCTAAC	AGACTTAGCA	TGGGTTTAGT	6000
	ATTTACATTC	ATACACCAAT	TCAATCATCT	CCTTTTTTTCA	TGTTTGAAGG	TTGATAGGTC	6060
	DITTUCTION	"""	TOTALIGUE CT	thication variable	TTCAACAAAT	CCACTGATGC	6120
	AGGAAATGTT	CATCACCAGT	TICAAAAGCI	TCTGACTGAA	1100000000	CCACIGAIGC	
75	ATATGAGCTG	AAGATCGCCA	ACAAGCTCTT	CGGAGAAAAG	ACGTATCAAT		6180
75						CTGAGTGGCC	
							6300
						CCAACGATAC	6360
							6420
	TONORAGE	TANTANTA	TARCATACCA	TCGATAAACT	GGCACTCACT	CAGTCACATA	6480
80	TOWOTAMONG	TOTAL TANGEN	TUNGWINCOW	**************************************	Publicay yang	GCVGCCLAMOR	6540
OU	CUATACATCT	1GG1GGGAAA	IGIATGACTA	AIGGGATATT	MATTOGWATEG		
	GTGAGTTCCT	GAGAATAGTT	GAGGAAGTAC	CAGGAAATAT	TGAATGCACA		6600
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	GAATCTCCTT	CAGGGAAGCC	TGCTCTGCAG	TTTGCAAACC	ACAGCCTCTT	CTGCTTCTGC	6720
_	CTTTTGCCAA	GATGATATTG	ACCTTCAGTG	ACCTCTTTCT	TGTGCCAGCC	CACATTCCCC	6780
85	تحليد لانكالياطيان	CCTACATCAC	ACCTGTATAA	AAATATCCAT	GGACAGGAGA	TACTGCATCT	6840
	TITIOCUTIO	TOTAL TOTAL	THE CHARMON	TACAAAAAA	TARCTTTCCT	AATATATAGT	6900
	ATTURGGETC	I GGATT CAGC	TINCIUTIGE	COMMONATA	***********	TOMACIACIAC	
	TACATAAATT	ACTCCTAATT	CCTACTTCTT	CCTTCATATC	TAAAGGAAT	ATTTAGATGC	6960

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CATCAAGAAA TTTTACCAGA CCAGTGTGGA ATCTACTGAT TTTGCAAATG CTCCAGAAGA 7020'
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                                                                            7080
       TTATAGAAC ACCTTGAGA AACCTATGCC AGTGAGCCTT GTGCTTGACA CTGCATGGGG
                                                                             7140
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                                                                             7560
                                                                             7620
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EEPCCNHPFL FFIRQNKTNS ILFYGRFSSP 5 Seq ID NO: 466 DNA sequence Nucleic Acid Accession #: NM_001910.1 Coding sequence: 50..1240 51 31 41 10 GGAGAGAAGA AAGGAGGGGG CAAGGGAGAA GCTGCTGGTC GGACTCACAA TGAAAACGCT 60 120 CCTTCTTTTG CTGCTGGTGC TCCTGGAGCT GGGAGAGGCC CAAGGATCCC TTCACAGGGT GCCCCTCAGG AGGCATCCGT CCCTCAAGAA GAAGCTGCGG GCACGGAGCC AGCTCTCTGA 180 GTTCTGGAAA TCCCATAATT TGGACATGAT CCAGTTCACC GAGTCCTGCT CAATGGACCA 240 GAGTGCCAAG GAACCCCTCA TCAACTACTT GGATATGGAA TACTTCGGCA CTATCTCCAT TGGCTCCCCA CCACAGAACT TCACTGTCAT CTTCGACACT GGCTCCTCCA ACCTCTGGGT 15 300 360 CCCCTCTGTG TACTGCACTA GCCCAGCCTG CAAGACGCAC AGCAGGTTCC AGCCTTCCCA 420 GTCCAGCACA TACAGCCAGC CAGGTCAATC TTTCTCCATT CAGTATGGAA CCGGGAGCTT 480 GTCCGGGATC ATTGGAGCCG ACCAAGTCTC TGTGGAAGGA CTAACCGTGG TTGGCCAGCA 540 20 GTTTGGAGAA AGTGTCACAG AGCCAGGCCA GACCTTTGTG GATGCAGAGT TTGATGGAAT 600 TCTGGGCCTG GGATACCCCT CCTTGGCTGT GGGAGGAGTG ACTCCAGTAT TTGACAACAT 660 GATGGCTCAG AACCTGGTGG ACTTGCCGAT GTTTTCTGTC TACATGAGCA GTAACCCAGA 720 780 AGGTGGTGCG GGGAGCGAGC TGATTTTTGG AGGCTACGAC CACTCCCATT TCTCTGGGAG CCTGAATTGG GTCCCAGTCA CCAAGCAAGC TTACTGGCAG ATTGCACTGG ATAACATCCA 840 25 GGTGGGAGGC ACTGTTATGT TCTGCTCCGA GGGCTGCCAG GCCATTGTGG ACACAGGGAC 900 TTCCCTCATC ACTGGCCCTT CCGACAAGAT TAAGCAGCTG CAAAACGCCA TTGGGGCAGC 960 CCCCGTGGAT GGAGAATATG CTGTGGAGTG TGCCAACCTT AACGTCATGC CGGATGTCAC 1020 CTTCACCATT AACGGAGTCC CCTATACCCT CAGCCCAACT GCCTACACCC TACTGGACTT 1080 CGTGGATGGA ATGCAGTTCT GCAGCAGTGG CTTTCAAGGA CTTGACATCC ACCCTCCAGC 1140 TGGGCCCCTC TGGATCCTGG GGGATGTCTT CATTCGACAG TTTTACTCAG TCTTTGACCG 30 1200 TGGGAATAAC CGTGTGGGAC TGGCCCCAGC AGTCCCCTAA GGAGGGGCCT TGTGTCTGTG CCTGCCTGTC TGACAGACCT TGAATATGTT AGGCTGGGGC ATTCTTTACA CCTACAAAAA 1320 GTTATTTTCC AGAGAATGTA GCTGTTTCCA GGGTTGCAAC TTGAATTAAG ACCAAACAGA 1380 ACATGAGAAT ACACACACA ACACACATAT ACACACACA ACACTTCACA CATACACACC 1440 ACTCCCACCA CCGTCATGAT GGAGGAATTA CGTTATACAT TCATATTTTG TATTGATTTT 35 1500 TGATTATGAA AATCAAAAAT TTTCACATTT GATTATGAAA ATCTCCAAAC ATATGCACAA 1560 GCAGAGATCA TGGTATAATA AATCCCTTTG CAACTCCACT CAGCCCTGAC AACCCATCCA 1620 CACACGGCCA GGCCTGTTTA TCTACACTGC TGCCCACTCC TCTCTCCAGC TCCACATGCT 1680 GTACCTGGAT CATTCTGAAG CAAATTCCGA GCATTACATC ATTTTGTCCA TAAATATTTC 1740 40 TAACATCCTT AAATATACAA TCGGAATTCA AGCATCTCCC ATTGTCCCAC AAATGTTTGG 1800 CTGTTTTTGT AGTTGGATTG TTTGTATTAG GATTCAAGCA AGGCCCATAT ATTGCATTTA 1860 TTTGAAATGT CTGTAAGTCT CTTTCCATCT ACAGAGTTTA GCACATTTGA ACGTTGCTGG 1920 TTGAAATCCC GAGGTGTCAT TTGACATGGT TCTCTGAACT TATCTTTCCT ATAAAATGGT 1980 2040 AGTTAGATCT GGAGGTCTGA TTTTGTGGCA AAAATACTTC CTAGGTGGTG CTGGGTACTT 45 CTTGTTGCAT CCTGTCAGGA GGCAGATAAT GCTGGTGCCT CTCTATTGGT AATGTTAAGA 2100 CTGCTGGGTG GGTTTGGAGT TCTTGGCTTT AATCATTCAT TACAAAGTTC AGCATTTT Seq ID NO: 467 Protein sequence Protein Accession #: NP_001901.1 50 21 31 MKTLLLLLV LLELGEAQGS LHRVPLRRHP SLKKKLRARS QLSEFWKSHN LDMIQFTESC SMDOSAKEPL INYLDMEYFG TISIGSPPQN FTVIFDTGSS NLWVPSVYCT SPACKTHSRF 120 QPSQSSTYSQ PGQSFSIQYG TGSLSGIIGA DQVSVEGLTV VGQQFGESVT EPGQTFVDAE 55 180 FDGILGLGYP SLAVGGVTPV FDNMMAQNLV DLPMFSVYMS SNPEGGAGSE LIFGGYDHSH 240 FSGSLNWVPV TKOAYWOIAL DNIQVGGTVM FCSEGCQAIV DTGTSLITGP SDKIKQLQNA 300 IGAAPVDGEY AVECANLNVM PDVTPTINGV PYTLSPTAYT LLDFVDGMQF CSSGFQGLDI 360 HPPAGPLWIL GDVFIRQFYS VFDRGNNRVG LAPAVP 60 Seq ID NO: 468 DNA sequence Nucleic Acid Accession #: NM_018058.1 Coding sequence: 319..1575 65 51 21 31 41 TACGCGCTGC GGGACCGGCA GGGGAACGCC ATCGGGGTCA CAGCCTGCGA CATCGACGGG GACGGCCGGG AGGAGATCTA CTTCCTCAAC ACCAATAATG CCTTCTCGGG GGTGGCCACG 120 TACACCGACA AGTTGTTCAA GTTCCGCAAT AACCGGTGGG AAGACATCCT GAGCGATGAG 180 GTCAACGTGG CCCGTGGTGT GGCCAGCCTC TTTGCCGGAC GCTCTGTGGC CTGTGTGGAC 70 AGAAAGGGCT CTGGACGCTA CTCTATCTAC ATTGCCAATT ACGCCTACGG TAATGTGGGC CCTGATGCCC TCATTGAAAT GGACCCTGAG GCCAGTGACC TCTCCCGGGG CATTCTGGCG 360 CTCAGAGATG TGGCTGCTGA GGCTGGGGTC AGCAAATATA CAGGGGGCCG AGGCGTCAGC 420 GTGGGCCCCA TCCTCAGCAG CAGTGCCTCG GATATCTTCT GCGACAATGA GAATGGGCCT 480 75 AACTTCCTTT TCCACAACCG GGGCGATGGC ACCTTTGTGG ACGCTGCGGC CAGTGCTGGT 540 GTGGACGACC CCCACCAGCA TGGGCGAGGT GTCGCCCTGG CTGACTTCAA CCGTGATGGC 600 660 AAAGTGGACA TOGTCTATGG CAACTGGAAT GGCCCCCACC GCCTCTATCT GCAAATGAGC ACCCATGGGA AGGTCCGCTT CCGGGACATC GCCTCACCCA AGTTCTCCAT GCCCTCCCCT 720 GTCCGCACGG TCATCACCGC CGACTTTGAC AATGACCAGG AGCTGGAGAT CTTCTTCAAC 780 80 AACATTGCCT ACCGCAGCTC CTCAGCCAAC CGCCTCTTCC GCGTCATCCG TAGAGAGCAC 840 GGAGACCCCC TCATCGAGGA GCTCAATCCC GGCGACGCCT TGGAGCCTGA GGGCCGGGGC 900 ACAGGGGTG TGGTGACCGA CTTCGACGGA GACGGGATGC TGGACCTCAT CTTGTCCCAT 960 GGAGAGTCCA TGGCTCAGCC GCTGTCCGTC TTCCGGGGCA ATCAGGGCTT CAACAACAAC 1020 TGGCTGCGAG TGGTGCCACG CACCCGGGTT GGGGCCTTTG CCAGGGGAGC TAAGGTCGTG 1080 85 CTCTACACCA AGAAGAGTGG GGCCCACCTG AGGATCATCG ACGGGGGCTC AGGCTACCTG 1140 TGTGAGATGG AGCCCGTGGC ACACTTTGGC CTGGGGAAGG ATGAAGCCAG CAGTGTGGAG 1200 GTGACGTGGC CAGATGGCAA GATGGTGAGC CGGAACGTGG CCAGCGGGGA GATGAACTCA 1260

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			AYDHSRVKLA				1800
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	VLAYYTVRNF	TLRNTKIKKG	SQKGRPSGRV	VTQYHYTQWP	DMGVPEYSLP	VLTPVRKAAY	1920
40			GTYIVLDSML				1980
40			LDSHIHAYVN				2040
			ERSRVGISSL				2100 2160
			I PDGQNMAED LEVRHFQCPK				2220
			HQLEKENSVD				2280
45						4-4	
43	-		ALPDGNIAES mence	LESLV			
	Seq ID NO: Nucleic Aci	574 DNA sec	ruence 1 #: Eos sec				
50	Seq ID NO: Nucleic Aci Coding sequ	574 DNA sec d Accession dence: 148-4	quence 1 #: Eos sec 1518	quence	41	51	
	Seq ID NO: Nucleic Aci	574 DNA sec d Accession	ruence 1 #: Eos sec		41 	5 1	
	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG	574 DNA sec d Accession dence: 148-4 11 CACGCACGAT	quence 1 #: Eos sec 1518 21 1 CTCACTTCGA	quence 31 TCTATACACT	 GGAGGATTAA	AACAAACAAA	60
50	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAAC	574 DNA sec. d Accession ence: 148-4	quence #: Eos sec 518 21 CTCACTTCGA CTCCCCCTCC	quence 31 TCTATACACT CTCTCCACTC	 GGAGGATTAA TGAGAAGCAG	AACAAACAAA AGGAGCCGCA	120
	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAAC CGGCGAGGGG	574 DNA sec. d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG	nuence 1 #: Eos sec 1518 21 CTCACTTCGA CTCCCCCTCC	quence 31 TCTATACACT CTCTCCACTC CGAATCCTAA	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT	AACAAACAAA AGGAGCCGCA CGCTTGCATT	120 180
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT	574 DNA second Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG	nuence 1 #: Eos sec 1518 21 CTCACTTCGA CTCCCCTCC TCTGGAAATG CCTGGATTGG	quence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA	120 180 240
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG	574 DNA sector dence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGCCTG	nuence 1 #: Eos sec 1518 21 CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG	120 180 240 300
50 55	Seq ID NO: Nucleic Ac: Coding sequ 1 	574 DNA sec. d. Accession nence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG	nuence #: Eos sec 518 21 CTCACTTCGA CTCCCCTCC CTGGAATGG CTCGGATTGG GTCCTATACA CCCAAAACAA	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA	120 180 240 300 360
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAA CAGCGAGGGG CAGCTCCTCT CTTGTTGAGA AAATATCCAA CAAGTAAATG	574 DNA sec. d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGCCTG CATGTAATAG TGAATCTTAA	nuence #: Eos sec 518	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT	 GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA	120 180 240 300
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAA CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA	574 DNA sec. d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC	nuence a #: Eos sec 518 21 CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCTGGATTGG GTCCTATACA CCCAAAACAA GAACTTAAA TGGGAAAACA	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TCTCCTATCA TTTCAGGGTT GTGGAAATTA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA	AACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT	120 180 240 300 360 420
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG	574 DNA sector accession a	nuence #: Eos sec 518	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TCTCCTATCA TCTCAGGGTT TTTCAGGAATTA AAAGCAACCA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TTGACTACCGT TCACTGGGGA	120 180 240 300 360 420 480 540 600
50 55	Seq ID NO: Nucleic Aci Coding sequ 1 	574 DNA sec d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGCCG CATGTAATAG TCATAACAC GAGTTTCAT TTCATAACAC TGTGTTCGAT TGTCATCTGA TCTCATCTGAT TCTCATCTGAT TCTCATCTGAT TCTCATCTGAT TCTCATCTGAT	quence #: Eos sec 518	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAG CGATTTTCAA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA	ACAAACAAA AGGAGCCGCA CGCTTGCATT CACAGGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTCAAA	120 180 240 300 360 420 480 540 600 660
50 55 60	Seq ID NO: Nucleic Aci Coding sequ CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA AAATATCCAA AACACATTCA GTCAGCGGAG AAATGCAAAT GAGATGCAAAA GGAAAAGGGA	574 DNA sec d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATAACAC GAGTTTCAGA TGTTATCATCTAT TCTATCTGCTT AGTTAAGACG	quence a #: Eos sec 518 21 } CTCACTTCGA CTCGCCCTCC TCTGGAATG GTCCTATACA CCCAAACAA AACGAAACAA AATGGTGTTT TGGATCAGA TGGGATCAGA TGATGCGGAC TTTATCCATT	Juence 31 TCTATACACT TCTATACACT CGAATCCTAA GCTAATGGAT TCTCCTATCA TCTCCTATCA TCTCCTATCA TCTCCAGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTACA TGTTTCAGG TTGTTTCAGG	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAC GTTTTGAGGA TTGGGACAGA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTTCCACTT AGCAGTCAAA AGAAAATTTG	120 180 240 300 360 420 480 540 600 660 720
50 55	Seq ID NO: Nucleic Aci Coding sequ 1	574 DNA sector accession a	quence #: Eos sec 5:18 21 CTCACTTCGA CTCCCCTCC CCTGGAAATG GTCCTATACA CCCAAAACAA GAAACTTAAA AATGGTGTTT TGGATCAGAG TGATGCGGAC TTTATCCATT TGGATCAGAA	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GGAGCACTGA TCTCCTATCA TCTCAGGGTT TTCAGGGTTA AAAGCAAGCA CATAGTTTAG CGATTTTCAA TTGTGGAATTTAA AAGTATTCAA AAGTATTAGAGAATTAA CATTTTCAGAGAATTTAA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGAGGA GTTTTGGGAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTCAAT AGCAGATCATT GCAGGCTGCT GCAGGCTGCT	120 180 240 300 360 420 480 540 600 660 720 780
50 55 60	Seq ID NO: Nucleic Ac: Coding sequ 1	574 DNA sector control of the contro	quence #: Eos sec 5:18 CTCACTTCGA CTCCCCTCC CTGGAATGG GTCCTATACA CCCAAACAA GAACTTAAA GAACTTAAA GATCGGATTT TGGATCAGAG TGTATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATT TGGATCCATA GAACCTTCTG	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTTA AAAGCAAGCA CATAGTTTAG CGATTTCAAG TTGTTTCAGA TTGTTTCAGA AGTGTTTAGAG AGTGTTAGTC CCAAACTCAA	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA TTGGGACAGA GTTTTGGGAAA CTTGACAAGTA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TCACTGCGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC	120 180 240 300 360 420 480 540 660 720 780 840
50 55 60	Seq ID NO: Nucleic Aci Coding sequ CACACATACG CAAAAAAAAC CAGCAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCAA GTCAGCGGAG AAATGCAATA GAGATGCAATA GAGATGCAATA GAGATGCAATA GAGATGCAATA AGGATACCAT AATGGCTCAT AATGGCTCAT	574 DNA sec. d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGCCG CATGTAATAG TCATAACAC GAGTTCAGA TGTCATCTGA TGTCATCTGA TCTACTGTT AGTTAAGAGC CGATTATGGT TAGTAACAC TCTACTGTT TGTCATCTGT TGTCATCTGT TGTCATCTGT TGACATCTCT TGACATCTCT TGACATCTCC	quence i #: Eos sec i= 1: Eos	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAGA TTGTTTGAG AGTGTTAGT CCAAACTCAA GACACAGTT	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCATCAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA TTGGGACAGA GTTTTGGGAA CTGACAAGTA ACTGACTGTTTT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCACTTGCAT TCACTGGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTTACATTTAC TTTTAAAGAT	120 180 240 300 360 420 480 540 660 720 780 840 900
50 55 60	Seq ID NO: Nucleic Acc Coding sequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG CAAGTAAATG CAAGTAAATG AACACATTCA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GAATGCAATA AGGATGCAAAT AGGATGAACA GGAAAAGGGA GATTCAAAG TTAGATCCAT AATGGCTCAT ACAGTTAGCA	574 DNA sec d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG CATGTAATAG TCATAACAC GAGTTCAGA TCTCATCTGA TCTCATCTGA TCTACTGT TCTACTGT TCATATGGT TCATATGGT TCATATGGT TCATATGGT TCATATGGT TCATATGT TCATATTGA TCATATCTGT TCATATCTGT TCATACTGT TGACATCTCC TCTCTGAAAG	puence ##: Eos sec 1518 21 CTCACTTCGA CTCCCCTCC TCTGGAATG CCTGGATTGG GTCCTATACA GAAACTAAA TGGGAAACA AATGGTGTTT TGGATCAGT TGGATCAGT TGATCCGGAC TTATCCATT TGGAGTCGAA GAACCTTCTG TCCCTGCACA CCAGTTGGCT	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTGAG CGATTTTCAG AGTGTTAGA AGTGTTAGA AGTGTTAGA AGTGTTAGA AGTGTTAGAG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTGTG	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAG TTTGAGGA TTGGGACAGA GTTTTGGGAA CTGACAAGTA ACTGATTTAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG ATCATTGGAA ATCATTGGAA ATCATTGGAA ATTCACTT AGCAGTCAAA AGGAAAATTTG GCAGGCTGCT TTACATTTAG TTATAAAGAT AATGCAACAA	120 180 240 300 360 420 480 540 660 720 780 840
50 55 60	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG GAATGCAATA GGAAAAGGGA GATTTCAAAG TTAGATCCAT AATGGCTCAT ACAGTTAGCA TCTGGTTATG TCTCTAGAG	574 DNA sec. d. Accession ence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG CAGTATAGGTTAATAG TCATACACC GAGTTCAGA TCTCATACAGA TCTACTCGAAAG CCGATTATGA TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC	puence #: Eos sec 518 21 CTCACTTCGA CTCCCCTCC CTGGAATGG GTCCTATACA CCCAAAACAA GAAACTTAAA GAGATCAGAG TGGATCAGAG TGGATCAGAG TGTATCACT TGGATCAGAG TGTATCCATT TGGATCAGAG GAACCTTCTG TCCCTGCACA CCAGTTGGCT GGACTACTTA GGACTACTTA GGACTACTTA GGACTACTTA GGACTACTTA GGACTACTTA GGACTACTTA GGACTACTTA GGACTACACT	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTAA AAAGCAAGCA CATAGTTTAG CGATTTTCAA TTGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACGTTG GTTTTTTTTG GTAAACAATT GGAAAGGAAG	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA AGATAACCTTA AGGACAAAACCTTT AAGGACAAAA TTGGGACAG GTTTTGGGAA CTGACAAGTA ACTGGATTGT AAGTTCTTG AAGTTCTTGAGAA ACTGGATTGT AAGTTCTTGAAAACCTT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA ATCATTGGGA ATTTCCACTT TCACTGGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGCAGTTTGT	120 180 240 300 360 420 480 540 660 720 780 840 900 960
50556065	Seq ID NO: Nucleic Aci Coding sequ CACACATACG CAAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCAA GTCAGCGGAG AAATGCAATA GGAAAAGGGA GATTTCAAAG TTTGATCAT AATGCTCAT AATGGCTCAT ACAGTTAGCA TCTGGTTTATG TTTCTAGAC AGTTCAGAAC AGTTCAGAAC	574 DNA sec. d Accession ence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGCCG CAGTAATAG TCATAACAC GAGTTCAGA TGTCATCAGA TGTCATCAGA TCTACTGTT AGTTAAGAGC CGATTATGA TCTACTGTT TGACATCTCT TCTCTGAAAG TCATGCTGT TCATGCTGT TCATGCTGTT TGACATCTCC TCTCTGAAAG TCATGCTGTT CAGAAAATGT CAGAAAAATGT CAGAAAAATGT CAGAAAAATGT	HENCE HE EOS SEC 1518 21 CTCACTTCGA CTCCCCCTCC TCTGGAATGG GTCCTATACA GCCAAAACAA GAACTTAAA TGGGAAACA AATGGTGTTT TGGATCAGAG TGATCCAGT TGGATCAGAG GAACCTTCTG GCACTCCTGCACA CCAGTTGGCT GGACTACACA CCAGTTGGCT TCCCTGCACA CTCATACACT TCAGGCTGAC TCAGGCTGAC TCAGGCTGAC	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CATAGTTTAGA CGATTTTCAA TGTTTGAGG AGTGTTAGTC CCAAACTCAA GACACAGTTG GTTTTTTTTTT	GGAGGATTAA TGAGAAGCA AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA TTGGGACAGA GTTTTGGGAA CTGACAAGTA ACTGACAGTA ACTGACAGTA ACTGACAGTA ACTGAGACCA ATACCAGCCT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG ATCATTGCAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTACATTTAC ATTTACATTAC TTACATTTAC TTACATTTAC TTACATTAC TTACATTAC TTTAAAGAT ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAACAA ACAGTACAAC TCTTGTTACA	120 180 240 300 360 420 480 540 660 720 780 840 900 960 1020
50556065	Seq ID NO: Nucleic Acc Coding sequ CACACATACG CAAAAAAAA CAGCGAGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG GAATGCAATA GGAAAAGGGA GAATTCAAAG TTAGATCAAT ACAGTTAAAG TTAGATCCAT ACTGGTTATG TCTCTAGAG TCTCTTAGAA TCTCTTAGAAC TCTCTTAGAAAC TCTGGTTATG TCTCTTAGAAC TCTGGTAAAAC TGGGAAAAGAC	574 DNA sec. d Accession tence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG GAGATTGCCG CAGTTAATAG TCATAACAC GAGTTCAGA TCTCATCTGA TCTACTGTT AGTTAAGAGC CGATTATGAT TCATCTGTT TGACATCTGT TCATCTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CAGAGTCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGAAAATGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT CTCAGATCGT	Hence HI: Eos sec 1518 21 CTCACTTCGA CTCCCCCTCC TCTGGAATG CCTATACA CCCAAACAA GAAACTTAAA TGGGATACA TGGATCGGA TTGATCGGAC TTATCCATT TGGATCGAG GAACCTTCTG TCCCTGCACA CCAGTTGGCT GGACTACTTA CTCATACACT TCAGGCTGAC TCAGCCTCC TCAGCCTCC TCAGCCTCC TCAGCCTCC TCAGCCTCC TCAGCCTCAC TCAGCCTCAC TTATGATACC TTATGATACC	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAAATTA AAAGCAAGCA CGATTTTCAG CGATTTCAG GGTGTTAGT CAAACTCAA CCACAGTTG CTTTTTTGTG CAAAACAATT GGAAAAGAAT CCAGAGAATT ATGATTGAG ATTTTTTTTTGTG CAAAACAATT ATGATTGAGAAATTAA CCAGAGAATT ATGATTGAGA	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACC ATCTCACTAA AGATAACTTT AAGGACAAA TTTGAGGA TTTGAGGA CTGACAAGT ACTGCACAGT ACTGCACAGT ACTGCACAGT AAGTTCTTAC AGATTCATAA AGATTCATAC AGATTCATAA AGATTCATAC AGATTCATAC AGATTCATAC AGATTCATGA	ACAAACAAA AGGAGCAGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG ATCATTGGAA ATCATTGGAA ATCATTGGAA AGCAGTCAAA AGGAGTCAAA AGGAGTCAAA AGAAATTTG CAGGCTGCT TTACATTTGA TTACATTTA AGAAAATTTG AATGCAACAA ACAGTACAA ACAGTACAAG AGCAGTTTTA TCTTGTTACA TTTGTACAG	120 180 240 300 360 420 480 540 600 720 780 840 900 960 1020
50556065	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGAGAG AAATGCAATA GAAAAGGGA GAAAAGGGA GAATGCAATA ATGGCTCAT ATGGTTCAT ATGGTTATG TCTGGTTATG TCTCTGGTATG TCTCTTGAGAC CAGTTCAGAAC CTGGGAAAGAC CCAGTTAGATC CGGGAAAGAC CCAGTTGGATG	574 DNA sec. cd Accession ence: 148-4 11	TLENCE TOTAL	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GGAGCACTGA TCTCCTATCA TTTCAGGGTT ATAGGAATTTAA AAAGCAAGCA CATAGTTTAG CGATTTCAA GACACAGTTA GACACAGTTA GTTTTTTTGTG CAAACAATT GGAAACAATT GGAAACAATT GGAAACAATT GGAAACAATT ATGATTGAGA ACAGTAT ATGATTGAGA ACAGTTT	GGAGGATTAA TGAGAAGCAG AGCGTTTCCT ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAACC ATCTCACTAA AGATAACTTT AAGGACAAAGTA GTTTTGAGGA GTTTTGAGGA GTTTTGGGAA ACTGGATTGT AAGTTCTTAC TCGAGAGCA AGATTCATGA AGATTCATGA AGATTCATGA ATACCAGCT AGTTTGCAGT CAGATGGCTA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAA ATTACATTTAA ACAGTACAA ACAGTACAA ACAGTACAA ACAGTACAA TCTTGTTACA TTTGTTACCAG TCTAGAGATTCCAGGTACAGA TCTTGTTACCAGGTACAGA TCAAGACTTG	120 180 240 300 360 420 480 540 600 660 720 780 840 960 1020 1080 1140 1200 1260
5055606570	Seq ID NO: Nucleic Ac: Coding sequ 1	574 DNA sec. d. Accession nence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TCATACAGA TGTCATCTGA TCTACTGGTT AGTTAACAGT TCTACTGCT TCTACTGCT TCATACTGT TGACATCTCC TCTTGAAAG TCATACTGTT TGACATCTCC TCTCTGAAAG TCATGCTGAT AGGTGTTTTC CAGAAAATGT CTCGAGTCGT GAGAGGACCA TCAATAATTT	HENCE ##: EOS SEC 1518 21 CTCACTTCGA CTCCCCCTCC TCTGGAAATG CCCGAAACAA GAAACTTAAA GAAACTTAAA GAAACTTAACA TGGGATACAG TGATGCGAC TTTATCCATT TGGATCGAA GAACCTTCTG TCCCTGCACA CCAGTTGGCT TGGACTACACT TCAGTCGAC TCATACACT TCAGTCGAC CCAGTTGCT TCAGTCGAC CCAGTTGCT TCAGTCGAC CCAGTTGCT TCAGTCACAC CCAGTTGCT TCAGTCACAC CCAGTTGCT TCAGTCACAC CCAGTTGCT TCAGTCACAC CCAGTTACACT TTATGATACC CAACCAAGCAT GCTACCAACCAT	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT GTGGAATTTAG CATAGTTTAG CATAGTTTAG CATAGTTTAG CATAGTTTAG GACACAGTT GCAAACTCAA GACACAGTT GGAAACAAT GCAAACAAT GCAAACAAT GGAAACAAT CAAACAAT CAAACAAT ATGATTAGAT ATGATTAGAT ATGATTTAGAT ATGATTTTAGA ATGAGTTATT	GAGGATTAA TGAGAAGCA AGCGTTTCCT ACTACAGACA ATCAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGAGGA CTGACAGA GTTTTGAGAA CTGACTTA ACTGATTTA ACTGATTTA ACTGATTTTA ACTGATTTTA ACTGATTGTA ACTTGACAGTA ACTTGACAGTA ACTTGACAGTA ACTTGATTTTA ACTTCACAGACCT AGATTGCACTA AGTTTGCAGAT AGTTTGCAGAT TTCTTCAGAT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAG AGGAGTTTGT TCTTGTTACA TTGTACCAG TCTTGTTACA TTGTACCAG TCTTGTACCAG TCAAGACTTG	120 180 240 300 360 420 480 540 660 720 840 900 900 1020 1140 1200 1260 1320
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505560657075	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAC CGGGGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGGAG AAATGCAATA GGAAAAGGGAATCCAT ACAGTTAGAC TTGGTTGAAG TTAGATCCAT ACAGTTAGCA TCTGGTATG TCTCTAGAC AGTTCAGAAC CGGAAAAGGGAAATCCAGATGGAAGAC CAGTTGGATG GGTGCTATTC GGCACTAATG AATCCTGAAC AAGGGGAAAAAGA AACCAAATCA AAGGGTGATG AACCAAATCA AAGGGTACTT GCACAAATCA AAGGGTACTT	574 DNA sec. d. Accession ence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG AGATTGGCTG CATGTAATAG TGAATCTTAA TTCATACAC GAGTTTCAGA TCATACTGTT AGTTAAGAG CGATTATTGA TCATACTGTT TGACATCTCT TCTGAAAG TCATGCTGAT TCATGCTGAT CTCGAGTCGT TCATGCAGTCT TCAGATCGT TCAGAGTCGT TCAGAGTCGT TCAGATCGT TCAGATCGT TGACATATTT GCTATATGG TCAGATCGT TGACATTTT GCTATATGG TCAGATCGT TTGACTTTT CAGATCGT TTGACTTTT CAGATCGAT AGGCATAATGA ATGAAGCCAA ATGAAGCCAA ATGAAGCCAA ATGAAGCCAA ATGAAGCCATTC CAGCCTCTTT CAGCCTCTTT	THENCE ##: EOS SEC 1 #: EOS SEC 1 #: EOS SEC 21 CTCACTTCGA CTCCCCTCC TCTGGAATG CCTGGATTGG GTCCTATACA CCCAAACAA GAAACTTAAA TGGGAAACA TGGGATCGGAC TTATCCATT TGGATCGGAC TCAGTTGGCT GGACTACT TCAGGTGAC CCAGTTGGCT TCAGGCTGAC TTATGATAC CTCATACACT TCAGGCTGAC TTATGATAC CTCATACACT TCAGGCTGAC AACCAAGCAT GCTACCCAAT AAAATACAGC CCCTGAATT AGAAGGCGT ACCCCAGATT GACTAACCGA ATCTTTAAAT GACTAACCGA ATCTTTTAAAT GACTAACCGA ATCTTTTAAG AAATGATGG GACTTCTCAG GACTTCTCAG AAATGATGGC	Juence 31 TCTATACACT CTCTCCACTC CGAATCCTAA GGAGCACTGA TCTCCTATCA TTTCAGGGTT TTGAGGAATTTA AAAGCAAGCA CATAGTTTAG CGATTTCAG GATTTCAG GACACTCAA GACACTCAA GACACTCAA GACACTCAA GACACTCAA GACACTCAA ATGATTTTTG GAAACAATT GGAAACAATT ATGATTCAG ATTGATTCAG ATTGATCACAA ATTGAACTG ATTGAACTC TCTACCACAA TCCACATCC ACTTCCC ACTTCCC ACTTCCC ACTTCCC ACTTCCC ACTTCCC TCTAAAACTG TCTAAAACTG	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA GTTTTGGGAA GTTTTGGGAA ACTGGATTGTA ACTGGATTGTA ACTGGATTGTA ACTGGATTGTA ACTGGATTGTA CTGAGAGCA TTCTCAGAT TTCTCAGAT TTCTCAGAT TTCTCAGAT TTCTCAGAT TTCTCAGAT TTCTCAGAT ACTGCACAC AGATGACTA AAGAAATAAT CTGGTAGAGA CACCACCAA AACTGCCACC TTCTTAGATC TTCTCAGAT AAGAAATAAT CAGCAGTCAA AACTGCCACC TTCTTAGATC	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTAAAA AGAAAATTTG GCAGGCTGCT TTACATTTAA AGAAAATTTG TTACATTTAA ACAGTACAA TCTGTTAC AGTAGCCATA GCCATACTGAT CAAGACTTG AAGAGAGGAG ATTCTCTGAA TCGCATAGGC TCACACTGTG TCACACTGTG TCACACTGTG TCACACTGTG	120 180 240 300 360 420 480 660 720 840 900 900 1020 1140 1260 1320 1380 1440 1560 1560 1620 1680 1740 1800
505560657075	Seq ID NO: Nucleic Ac: Coding sequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GAATGCAATA GAATGCAATA GGAAAAGGGA GATTCCAAAG TTAGGATCAT ACTGGTTATG TCTCTAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGTTCAGAC AGGTACAT AAGGAAAAAG AACCAAATCA AACGAAAAAC AAGGTGATG AACGAAAACA AACGAAATACA AAGGGTGATG AACGAAAATACA AAGGGTGATG ACAGAAAAAC AAGGAGAAAAAC AAGGAGAAATACA AAGGAAAATACA AAGGAGAACTT AACTTGTCGG	574 DNA sec. d Accession ence: 148-4 11 CACGCACGAT ATTTCCTTCG CCGCAGACCG GTGTTTGCCG GAGTTTGCCG GAGTTAATAG TCATACAGA TCTCATACAGA TCTATACAGA TCTACTGCT AGTTAAGAGC CGATTATGA TCATACTGTT TGACATCTC TCTCTGAAAG TCATGCTGAT TCAGATCTC TCTGAGATGT TCAGATCTT GAGAGACCA TCATATATTCCT TGAGACCT TCAGATATTT GCTTATATGG TTGATCTTT AGACATTTT AGACATTTT AGACATTTT CCTATATTGA TCATATTCCT TGAAAAAGGA ATGAACCAA TTCCCAATAC ATATTTCCTT CAGCCTCTTT CAGCTCTCTT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTTT CAGCCTCTT CAGCCTCT CAGCCTCT CAGCCTCT CAGCCT CAGCC CAGCCT CAGCC CAGCCT C	Hence #: Eos sec 1 #: Eos sec 1 #: Eos sec 1 #: Eos sec 21	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT GGAGCACTGA TCTCCTATCA TTTCAGGGTT GTGGAATTTAG CATATTGAGG CATAGTTAGA CATAGTTTAGA GACACAGTT GGAACCAAA GACACAGTT GGAAACTAAA GACACAGTT GGAAACTAAT GGAAAGAAT ATGAATTTGAA GAATTTTGA GAAACTAAT ATGAATTTGA GAAACTAAT TCGAACTGA ATTGGAACT ACTGTGACT ACCACTTCC ACTGTGACT ACAGTTTCTA	GGAGGATTAA TGAGAAGCA ACCAAAAAAA ATATTGATGA GGGATAAAAC ATCTCACTAA AGATAACTTT AAGGACAAAA GTTTTGAGGA TTGGGACAGA CTGACAGAT ACTGACAGTA ACTGACAGTA ACTGACAGTA ACTGCACTAC AGATGACTTT AGGATGATT TTGAGAGCA TTGTGAGAGCA ATACCAGCCT AGTTTGCAGAT TTGTCGAAT TTGTCGAAT TTGTCGACAT TTGTCGACAT TTGTCGACAT AGAATAAT TTGTCGACAT AGAATGAT AGAATGAT AGAATGACA CACACTACAA CACACTACAAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG ATCATTGCAA TGACTACCGT TCACTGGGA ATTTCCACTT AGCAGTCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAC TTTTAAAGAT AATGCAACAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACAAA ACAGTACCAG TCATTGTACCAG TCATGTACA TCTGTTACA TTGTACCAG TCAAGACTTG CCAGGCGCTAC AGTACCAG AGTACCAG AGTACCAG ATTCTCTGGA ACAGCTCTGAT CAAGACTTC CAAGACTTC CAAGACTTC CAAGACTTC TCACACTGGT TCACACTGTGT TCACACTGTGT TCACACTGTGT TCACACTGTGT TCACACTGTGT TCACACTGTGT TCACACTGTGT TCACACTACTGCT TCACACTACTC TCACACTACTC TCACACTACT TGAGGGAGGAG	120 180 240 360 420 480 660 720 780 840 960 1020 1140 1260 1320 1380 1440 1560 1680 1740 1680 1740 1800
50 55 60 65 70 75	Seq ID NO: Nucleic Aci Coding sequ 1 CACACATACG CAAAAAAAC CGGCAGGGG GAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATA GAGATGCAATA GAGATGCAATA GAGATGCAAT AATGCATA AATGCTAT ACAGTTAGCA TTCTCTAGAC TTCTCTAGAC TTCTCTAGAC ACTTCAGAC ACGAAATCA AACGAATACA AACGAATTAA ACGAAATCA ACGAATTACA ACGAATCA ACTTCTCTCA ACTTCTCA ACTTCTCTCA ACTT	574 DNA sec. d. Accession ence: 148-4 11	quence #: Eos sec 518 21 CTCACTTCGA CTCCCCTCC CTGGAAATG GTCCTATACA CCCAAAACAA GAAACTTAAA AATGGTGTTT TGGATCGGA TTTATCCATT TGGATCGGA GAACCTTCTG GACTACTCA CCAGAAACAA CAGTTGGCT TCAGTCGAT TCAGTCGAT TCATACACT TCAGTCGCT CCAGTTGCC CGACTACTTA CTCATACACT CTCATACCTA AAATACAGC CCCTGGATTA AAATACAGC CCCTGGATTA AGAAGGCGCT ACCCAGATTT GACTAACCGA AAATGATGGC CATCCTTAAAT GACTTCTCAG CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGATACT CCT	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GGAGCACTGA TCTCCTATCA TTTCAGGGTT TTCAGGGTT AAAGCAAGCA CATAGTTTAG CGATTTTCAG AGTTTTGAG AGTGTTAGTC CCAAACTCAA GACACAGTTG GAAACAATT GGAAACAATT GGAAACAATT GGAAACAATT GAAACAATT GAAACAATT GAACACTGA ATTGTGAGT ATTGTGACTC ATTGTGAACT ATTGTGAACT ATTGTGAACT ATCCCCAACAA TCCACCTCCC ACTGTGACTG GCAGGTTCT CTAAAACTG ACTGTGACTG ACAGTTCTCA GCAGCTGAAACTG ACAGTTCTCA GCAGCTGAAACTG ACAGTTCTCA GCAGCTGAAACTG ACAGTTCTCA GCAGCTGAAACTG ACAGTTCTCAAAACTG ACAGTTCTCAAAACTG ACAGTTCTCAAAACTG GCAGCTGAAAC	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAACC ATCTACTAA AGATAACTT AAGGACAA ATATTGATGA GGATAAACT ATGGACAGA TTGGGACAGA GTTTTGAGGA CTGACAAGTA AAGTTCTTAC TTCGAGAGCA AGATTCATGA AGAATTAAT CTGGTAGGA AACAGCAT AAGAAATAAT CTGGTAGGA AACAGTCAC AACTGCCAC TTCTTAGATC TTCTTAGATC TTAACAGAATA AATCAGAATA AATTCTTCAGG	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAA TTGGGGAAAA TGACTACCGT TCACTGGGGA ATTTTCACTT AGGAGTCTAAA AGGAGTCTAAA AGGAGTCTAA AGGAGTCAAA AGGAGTCAT TTATAAAGAT TCATTTGACAT TCTTGTACAG TCTGTACAG TCTAGAGACTTG AGTAGCATA AGCAGTTCTA CAAGACTTG AGTAGCATA CCTACTGAT CCAAGACTTG TCAAGACTTG TCTTGTACA TTTGTACCAG TCAAGACTTG TCTTGTACCAG TCAAGACTTG TCACACTTTG TCACACTTTG TCACACTTTG TCACACTTCT TCACAGTCCC	120 180 240 300 360 420 480 660 720 780 960 1080 1140 1200 1320 1380 1440 1560 1680 1740 1800 1900
505560657075	Seq ID NO: Nucleic Aci Coding sequ 1	574 DNA sec cd Accession ence: 148-4 11	THENCE ##: EOS SEC 1 #: EOS SEC 1 #: EOS SEC 1	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GGAGCACTGA TCTCCTATCA TTTCAGGGTT ATGGAATTTAG CGATTTCAG CGATTTCAG CGATTTCAG CGATTTCAG CGATTTCAG GCTATTCAG ATGGTTAGT CCAAACTCAA GACACAGTTG GAAACAATT GGAAAGAAT GGAATTTTGG GATTTTTGG CAAACAAT ATGGATTTGA ATGGATTAG CCAGAGAAT ATGGAACTGA ATTGGAACT ATGGAACT ATTGGAACT ATTGGAACT ATTGGAACT ATTGGAACT ATTGGAACT ATTGCACAA ATCCCCAACAA TCCCCAACAA TCCACTTCC ACTGTGCT CTAAAACT ACAGTTTCTA ACAGTTACCC ACTGTGACG ACATTACCC ACTGTGACG ACATTACCC ACTGTGACG ACATTACCC	GGAGGATTAA TGAGAAGCA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCTACTAA AGGATAAAC ATCTCACTAA AGGATAAAC ATCTCACTAA AGGATAAAC ATCTCACTAA AGGACAAAA ATATTGAGGA TTGGGGACAGA GTTTTGAGGA GTTTTGAGAGTA AAGTTCTTAC TTCGAGAGCA ATACCAGCT AGATTCTTAC TTCTCAGAT TTGTCGACAT TTGTCGACAT TTGTCGACAT AGGATTACAA CAGATGACAA ACAGCTA AAGAAATAAT CTGGTAGAG CACACTACAA AAGGAATTC AACTGCAC TTCTTAGATC TAACAGACT AACTGCACC TTCTTAGATC TAACAGAATAA ATTCTTCAGG AAGGGTATAT	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTCAAA AGGAGCTGCT TTACATTTAAAGAT AATGCACTACATTTAAAGAT ACAGTACAAG ACAGTACCAG TCAAGACTTG TACATTTGTTACA TCTTGTTACA TCTTGTTACA TCTTGTTACA TCTTGTTACA TCTTGTTACA TCTTACTACAGT CCACACTTG TCACACTTG TCACACTTG TCACACTTG TCACACTTG TCACACTTG TCACACTTCC ATTTTCCTCC	120 180 240 300 360 420 660 720 840 900 1020 1080 1140 1260 1320 1380 1560 1680 1740 1800 1800 1900
50 55 60 65 70 75	Seq ID NO: Nucleic Aci Coding Bequ 1 CACACATACG CAAAAAAAC CGGCGAGGGG CAGCTCCTCT CTTGTTGAAG AAATATCCAA CAAGTAAATG AACACATTCA GTCAGCGAGG AAATGCAATA GGAAAAGGAGTACATA ACAGTTAGAT TTAGATCCAT TAGATCCAT ACAGTTAGCA TCTGGTATG TCTCTAGAC TCTGTATG TCTCTAGAC CAGTTAGGAT CAGTTAGAT GGAAAAGC GAAGAGGGAA AACCAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAATCA ACGAAAAAG GAAGGTACTT ACTTGTCGG ACTTTTTTG GAAAACCTTCTG GAAAACCCAG	574 DNA sec. d. Accession lence: 148-4 lance: 148-4 lance	quence #: Eos sec 518 21 CTCACTTCGA CTCCCCTCC CTGGAAATG GTCCTATACA CCCAAAACAA GAAACTTAAA AATGGTGTTT TGGATCGGA TTTATCCATT TGGATCGGA GAACCTTCTG GACTACTCA CCAGAAACAA CAGTTGGCT TCAGTCGAT TCAGTCGAT TCATACACT TCAGTCGCT CCAGTTGCC CGACTACTTA CTCATACACT CTCATACCTA AAATACAGC CCCTGGATTA AAATACAGC CCCTGGATTA AGAAGGCGCT ACCCAGATTT GACTAACCGA AAATGATGGC CATCCTTAAAT GACTTCTCAG CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGAATAC CATCCTTAAAT CCTTGAATAC CCTTGATACT CCT	JUENCE 31 TCTATACACT CTCTCCACTC CGAATCCTAA GCTAATGGAT TCTCCTATCA TTTCAGGGTT AGAGCAGCA CATAGTTTAG CGATTTCAG GATTTCAG GATTTTCAG GATTTTCAG GACACGTA GCAAACTCAA GACACGTAG CAAACAATT GGAAACAATT GGAAACAATT GGAAACAATT CAAAACAATT ATGATTGAG ATTGGAACTA ATGATTTGAACTG ATTGGAACTA TCCACACAA TCCACTACAA TCCACTTCCC TCTAAAACTG ACAGTTCCA CAGGTTTCTA CAGATATCCC CTTATACCAC CTTATACCAC	GGAGGATTAA TGAGAAGCA ACTACAGACA ATCAAAAAAA ATATTGATGA GGGATAAAC ATCTCACTAA AGATAACTT AAGGACAAAA AGATTACACTA AGATAACTT AAGGACAAAA GTTTTGAGGA GTTTTGAGGA GTTTTGGGAA ACTGCACAGA AGATTCATAC AGATTCATAC AGATTCATAC AGATTCATAC AGATTCATCA TTCTCAGAT TTCTCAGAT TTGTCGACAT AGATACAGCCT AGATACAGCCT ACAGAGAAAAAAAAAA	ACAAACAAA AGGAGCCGCA CGCTTGCATT ACAGAGAAAA TTGGGGAAAG AGATCTTACA ATCATTGGAA TGACTACCGT TCACTGGGGA ATTTCCACTT AGCAGTAAAA AGGAGATCAAA AGAAAATTTG GCAGGCTGCT TTACATTTAA AGAAAATTTG TTACATTTAA ACAGTACAAG AGCAGTACAAG AGCAGTACAAG AGCAGTACAAG AGCAGTACAAG AGCAGTACAAG AGCAGTACAAG ACAGTACCAG TCAAGACTTG AGTAGCCATA CCAGTCCAGT	120 180 240 300 360 420 480 660 720 780 960 1080 1140 1200 1320 1380 1440 1560 1680 1740 1800 1900

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	CTAAAATTAG	GCAGGAAAAT	CAGTCTAGTT	CTGTTATCTG	TTGATTTCCC	ATCACCTGAC	7200
00	AGTAACTTTC	ATGACATAGG	ATTCTGCCGC	CAAATTTATA	TCATTAACAA	TGTGTGCCTT	7260
80	TTTGCAAGAC	TTGTAATTTA	CTTATTATGT	TTGAACTAAA	ATGATTGAAT		7320
	TTCTAAGAAT	GGAATTGTGG	TATTTTTTC	TGTATTGATT	TTAACAGAAA		7380
	ATAGAGGTTA	GGAATTCCAA	ACTACAGAAA	ATGTTTGTTT	TTAGTGTCAA	ATTTTTAGCT	7440
	GTATTTGTAG	CAATTATCAG	GTTTGCTAGA	AATATAACTT	TTAATACAGT	AGCCTGTAAA	7500
0.5	TAAAACACTC	TTCCATATGA	TATTCAACAT	TTTACAACTG	CAGTATTCAC	CTAAAGTAGA	7560
85	AATAATCTGT	TACTTATTGT	AAATACTGCC	CTAGTGTCTC	CATGGACCAA	ATTTATATTT	7620
	ATAATTGTAG	ATTTTTATAT	TTTACTACTG	AGTCAAGTTT	TCTAGTTCTG	TGTAATTGTT	7680
	TAGTTTAATG	ACGTAGTTCA	TTAGCTGGTC	TTACTCTACC	AGTTTTCTGA	CATTGTATTG	7740

TGTTACCTAA GTCATTAACT TTGTTTCAGC ATGTAATTTT AACTTTTGTG GÁAAATAGAA 7800 ATACCTTCAT TTTGAAAGAA GTTTTTATGA GAATAACACC TTACCAAACA TTGTTCAAAT 7860 Α ΑΑΑΑΑΑΑΑ ΑΑΑΑΑΑΑΑΑ Seg ID NO: 583 Protein sequence Protein Accession #: NP_002842.1 10 MRILKRFLAC IQLLCVCRLD WANGYYRQQR KLVEEIGWSY TGALNQKNWG KKYPTCNSPK QSPINIDEDL TOVNVNLKKL KFQGWDKTSL ENTFIHNTGK TVEINLTNDY RVSGGVSEMV 120 PKASKITFHW GKCNMSSDGS EHSLEGQKFP LEMQIYCFDA DRFSSFEEAV KGKGKLRALS 180 ILFEVGTEEN LDFKAIIDGV ESVSRFGKQA ALDPFILLNL LPNSTDKYYI YNGSLTSPPC 240 15 TDTVDWIVFK DTVSISESQL AVFCEVLTMQ QSGYVMLMDY LQNNFREQQY KFSRQVFSSY 300 TGKEEIHEAV CSSEPENVQA DPENYTSLLV TWERPRVVYD TMIEKFAVLY QQLDGEDQTK 360 HEFLTDGYQD LGAILNNLLP NMSYVLQIVA ICTNGLYGKY SDQLIVDMPT DNPELDLFPE 420 LIGTEEIIKE EEEGKDIEEG AIVNPGRDSA TNQIRKKEPQ ISTTTHYNRI GTKYNEAKTN 480 RSPTRGSEFS GKGDVPNTSL NSTSQPVTKL ATEKDISLTS QTVTELPPHT VEGTSASLND 540 20 GSKTVLRSPH MNLSGTAESL NTVSITEYEE ESLLTSFKLD TGAEDSSGSS PATSAIPFIS 600 ENISQGYIFS SENPETITYD VLIPESARNA SEDSTSSGSE ESLKDPSMEG NVWFPSSTDI 660 TAQPDVGSGR ESFLQTNYTE IRVDESEKTT KSFSAGPVMS QGPSVTDLEM PHYSTFAYFP 720 TEVTPHAFTP SSRQQDLVST VNVVYSQTTQ PVYNGETPLQ PSYSSEVFPL VTPLLLDNQI 780 LNTTPAASSS DSALHATPVF PSVDVSFESI LSSYDGAPLL PFSSASFSSE LFRHLHTVSQ 840 25 ILPQVTSATE SDKVPLHASL FVAGGDLLLE PSLAQYSDVL STTHAASETL EFGSESGVLY 900 KTLMFSQVEP PSSDAMMHAR SSGPEPSYAL SDNEGSQHIF TVSYSSAIPV HDSVGVTYQG 960 SLFSGPSHIP IPKSSLITPT ASLLQPTHAL SGDGEWSGAS SDSEFLLPDT DGLTALNISS 1020 PVSVAEFTYT TSVFGDDNKA LSKSEIIYGN ETELQIPSFN EMVYPSESTV MPNMYDNVNK 1080 LNASLOETSV SISSTKGMPP GSLAHTTTKV FDHEISQVPE NNFSVQPTHT VSQASGDTSL 1140 30 KPVLSANSEP ASSDPASSEM LSPSTQLLFY ETSASFSTEV LLQPSFQASD VDTLLKTVLP 1200 AVPSDPILVE TPKVDKISST MLHLIVSNSA SSENMLHSTS VPVFDVSPTS HMHSASLQGL 1260 TISYASEKYE PVLLKSESSH QVVPSLYSND ELFQTANLEI NQAHPPKGRH VFATPVLSID 1320 EPLNTLINKL IHSDEILTST KSSVTGKVFA GIPTVASDTF VSTDHSVPIG NGHVAITAVS PHRDGSVTST KLLFPSKATS ELSHSAKSDA GLVGGGEDGD TDDDGDDDDD RDSDGLSIHK 1440 35 CMSCSSYRES QEKVMNDSDT HENSLMDQNN PISYSLSENS EEDNRVTSVS SDSQTGMDRS PGKSPSANGL SQKHNDGKEE NDIQTGSALL PLSPESKAWA VLTSDEESGS GQGTSDSLNE 1560 NETSTDFSFA DTNEKDADGI LAAGDSEITP GFPQSPTSSV TSENSEVFHV SEAEASNSSH 1620 ESRIGLAEGL ESEKKAVIPL VIVSALTFIC LVVLVGILIY WRKCFQTAHF YLEDSTSPRV 1680 ISTPPTPIFP ISDDVGAIPI KHFPKHVADL HASSGFTEEF ETLKEFYQEV QSCTVDLGIT 1740 40 ADSSNHPDNK HKNRYINIVA YDHSRVKLAQ LAEKDGKLTD YINANYVDGY NRPKAYIAAQ 1800 GPLKSTAEDF WRMIWEHNVE VIVMITNLVE KGRRKCDQYW PADGSEEYGN FLVTQKSVQV 1860 LAYYTVRNFT LRNTKIKKGS QKGRPSGRVV TQYHYTQWPD MGVPEYSLPV LTFVRKAAYA 1920 KRHAVGPVVV HCSAGVGRTG TYIVLDSMLQ QIQHEGTVNI FGFLKHIRSQ RNYLVQTEEQ 1980 YVFIHDTLVE AILSKETEVL DSHIHAYVNA LLIPGPAGKT KLEKQFQLLS QSNIQQSDYS 2040 45 AALKQCNREK NRTSSIIPVE RSRVGISSLS GEGTDYINAS YIMGYYQSNE FIITQHPLLH 2100 TIKDFWRMIW DHNAQLVVMI PDGQNMAEDE FVYWPNKDEP INCESFKVTL MAEEHKCLSN 2160 EEKLIIQDFI LEATQDDYVL EVRHFQCPKW PNPDSPISKT FELISVIKEE AANRDGPMIV 2220 HDEHGGVTAG TFCALTTLMH QLEKENSVDV YQVAKMINLM RPGVFADIEQ YQFLYKVILS 2280 LVSTRQEENP STSLDSNGAA LPDGNIAESL ESLV 50 Seg ID NO: 584 DNA seguence 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GCTGGGCCTC CTCCTGACGG AAATCGTGCG GTCTTGGTCG CTTGCACTGA 840 CTTGGGCATT GAATTACCGA ACCGGTGTCC GCTTGCGGGG GGCCATCCTA ACCATGGCAT TTAAGAAGAT CCTTAAGTTA AAGAACATTA AAGAGAAATC CCTGGGTGAG CTCATCAACA TTTGCTCCAA CGATGGGCAG AGAATGTTTG AGGCAGCAGC CGTTGGCAGC CTGCTGGCTG GAGGACCCCT TETTGCCATC TTAGGCATGA TTTATAATGT AATTATTCTG GGACCAACAG GCTTCCTGGG ATCAGCTGTT TTTATCCTCT TTTACCCAGC AATGATGTTT GCATCACGGC 1080 75 TCACAGCATA TTTCAGGAGA AAATGCGTGG CCGCCACGGA TGAACGTGTC CAGAAGATGA 1200 ATGAAGTTCT TACTTACATT AAATTTATCA AAATGTATGC CTGGGTCAAA GCATTTTCTC 1260 AGAGTGTTCA AAAAATCCGC GAGGAGGAGC GTCGGATATT GGAAAAAGCC GGGTACTTCC 1320 AGGGTATCAC TGTGGGTGTG GCTCCCATTG TGGTGGTGAT TGCCAGCGTG GTGACCTTCT 1380 80 CTGTTCATAT GACCCTGGGC TTCGATCTGA CAGCAGCACA GGCTTTCACA GTGGTGACAG 1440 TCTTCAATTC CATGACTTTT GCTTTGAAAG TAACACCGTT TTCAGTAAAG TCCCTCTCAG 1500 AAGCCTCAGT GGCTGTTGAC AGATTTAAGA GTTTGTTTCT AATGGAAGAG GTTCACATGA 1560 TARAGARCAR ACCAGCCAGT CCTCACATCA AGATAGAGAT GARARATGCC ACCTTGGCAT 1620 GGGACTCCTC CCACTCCAGT ATCCAGAACT 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16		586 DNA so id Accession uence: 89	n_#: NM_001	1327.1			
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40		588 DNA se id Accession uence: 524	ı#: Eos se	equence			÷
45	1	11 1	21 	31 1	41 	51 	
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505560	CTTCGTGGGC GAAGGCCAGG CCTGATGGCC GGTCCCCGGG CCGCATGGCG ATCAGCTCCT GTGTTTTTGG TAGGTCATGC GCCTGATTGT CTGAGCTA Seq ID NO: Protein Acc MQAEGQGTGG PRGPHCGAAS FLPVFLAQAP Seq ID NO: Nucleic Acc	CCTGACCTTC GCACAGGGGG CAGGGGGCAA GCGCAGGGGC GTGCCGCTTC TGCTTCCAGCT GTCTCCAGCT TTGTCGCTGG 589 Protes cession #: 1 1 1 STGDADGPGG AQDGRCPCGA	TCTCTGAGAG TCTCTGAGAG TCTGACGGCC AGCAAGGGCC TGCGCAGGAT CCGACTGACT CCTCAGGGCAG AGGGAATGGT AGGGAATGGT AGGAGACGG in sequence bos sequence PGIPDGPGGN RRPDSRLLQF equence 1 #: NM_0055	CCGGGCAGAG GATGCTGATG CCAGGAGAGG TCGGGGCCGA GGAAGGTGCC TTGATGTGAACC TTGATGTGAA AGGCGCTAAG CCCAGCACGA CTTACATGTT 31 AGGPGEAGAT RLTAADHRQL	 GCTCCGGAGC GCCCAGGAGG GCGGTGCCAC GAGGAGGCGC CCTGCGGGGC ACCGCCAACT TCACGCAGT TCACGCAGT TGTGCCAGT TGTTTCTGTA	CATGCAGGCC CCTGGCAT GGGCGGCAGA CCCGCGGGGT CAGGAGCTCTCC CTTTCTGCCC CGCCCCTTCC CATTGTGGGG GAAAATAAAG	120 180 240 300 360 420 540 600
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50556065	CCTCGTGGGC GAAGGCCAGG CCGATGGCC GGTCCCCGGG CCGCATGGCG GACAGCCGC ATCAGCTCATGC GTGTTTTTGG TAGGTCATGC CTGAGTTA Seq ID NO: Protein Act MQAEGQGTGG PRGPHGGAAS FLPVFLAQAP Seq ID NO: Nucleic Ac: Coding sequence ACAGCGGAGC AGACAGAGC GCTTCTCGCT ATGGGAAGTC	CCTGACCTTC GCACAGGGGG GCACAGGGGGC TGCTCAGGTT TGTCTCCAGCA CTCAGGCTCC TTGTCGCTGG 589 Protes CCCCTCTCCCCT TTGTCGCTGG 589 Protes CCCCTCTCCCCT 11 GCACAGGGGCCC CCCCCCCCCC 11 GCAGAGTGAG CCCCCCCCCCCCCCCCCCCCCCCCCCCC	TCTCTGAGAG TCTCTGAGAG TCTGACGGCC AGCAAGGGCC TGCGCAGGAT CCGACTGACT CCTCAGGGCAG AGGGAATGGT AGGAGATGGT AGGAGACGG in sequence by sequence pgippggga RRPDSRLLQF equence the NM_0055 3671 21 AACCACCAAC GGCACCGCCA ACCAGCCCGGG ATCTTGATC	CCGGGCAGAG GATGCTGATG CCAGGAGAGG TCGGGGCCGA GGAAGGTGCC GCTGCAGACC TTGATGTGGA AGGCGCTAAG CCCAGCACGA CTTACATGTT 31 AGGPGEAGAT ### AGGPGEAGAT	GCTCCGGAGC GCCCAGGAGG GCCCAGGAGG GCCCAGGAGG CCTGCGGGGC ACCGCCAACT TCACGCAGTG CCCAGCCTGG GTGGCCAGTT TGTTTCTGTA 41 GGGGPRGAGA QLSISSCLQQ 41 GGCAGCGACC CTGGCTGGGC CAGGCGACC CTGGCTGGGC CAGACAACT CAGACAAACT CAGACAAACT	CATGCAGGCC CCCTGGCATT GGGCGGCAGA CCCGCGGGGT CAGGAGGCCG GCAGCTCCC CATTGTGGGG GAAAATAAAG 51 ARASGPRGGA LSLLMWITQC 51 CCTGCAGCGG TGCTGCCTCT CTGCAGCGG TGCTGCTCT CATGCAGCGG TGCTGCAGCAG GGTAATGAAT	120 180 240 360 420 480 540 600 120
5055606570	CCTCGTGGGC GAAGGCCAGG CCGATGGCG GGTCCCCGGG CCGCATGGCG GACAGCCGC ATCAGCTCCT GTGTTTTTGG TAGGTCATGC CCGTGATGT CTGAGCTA Seq ID NO: Protein Acc MQAEGQGTGG PRGPHGGAAS FLPVFLAQAP Seq ID NO: Nucleic Acc Coding sequ ACAGCGGAGC AGACAGAGAC GCTTCTCGCT ATGGGAAGT TCCGCTGCCT GCTTTTACCG	CCTGACCTTC GCACAGGGGG GCACAGGGGG CAGGGGGCAA GCGCAGGGGC TGCTTCAGGTT TGTCTCCAGCA CTCAGGCTCC CTCTCCCCT TTGTCGCTG 589 Protei CCESSION #: I STGDADGPGG AQDGRCPCGA SGQRR 590 DNA Sc id Accession uence: 90 11 GCAGAGTGAG TGAGCGGCCC CAGGCAGTGA CACAGAGAAA	TCTCTGAGAG TTCGACGGGC TGCGCAGGGC AGCAAGGGCC TGCGCAGGAT CCGACTGACT CCTCAGGGCAG AGGGAATGGT AGGAAGGACG In sequence 21 PGIPDGPGGN RRPDSRLLQF equence 1 #: NM_0055 8671 21 AACCACCAAC GGCACCGCA GCAGCCCGCA GCAGCCCGGC AGGGACCGCT AGGGACCCGCT AGGGACCCGCT AGGGACCCGCT	CCGGGCAGAG GATGCTGATG CCAGGAGAGG GCAGAGGTGCC GCTGCAGACC TTGATGTGGA AGGCGCTAAG CCCAGCACGA CTTACATGTT 31 AGGPGEAGAT AGGCGCTAAG CCACTCAGA CCCAGCACGA CTTACATGTT CCAGGAGAGT CCCACTCAGG GCGAACTTCA GTTTGCCCTG	GCTCCGGAGC GCCCAGGAGG GCCCAGGAGG CCGGGGGC CCTGCGGGGC CCCAGCCAGCT CCCAGCCTGG GTGGCCAGT TGTTTCTGTA 41 GGRGPRGAGA QLSISSCLQQ 41 GGCAGCGACC CTGGCTGGG GAGGAAGTC CCGGAGAAGC CCTGGCTGGGC CCTGGCTGGC CCGCTGGC CCTGGCTGG	CATGCAGGCC CCCTGGCATT GGGCGGCAGA CCCGCGGGGT CAGGAGGCCG CATTCTCCCC CGCCCCTTCC CATTGTGGG GAAAATAAAG 51 ARASGPRGGA LSLLMWITQC 51 CCTGCAGCGG TGCTGCTCT TGTGATTGCA TGGAATGGAT TGCAAGAATG	120 180 240 360 420 480 540 600 120
505560657075	CTTCGTGGGC GAAGGCCAGG GCAGGCCGC GGTCCCCGGG CCGCATGGCG GGCCCCATGGCG GACAGCCGCC ATCAGCTCCT GTGTTTTTGG TAGGTCATGC GCCTGATTGT CTGAGCTA Seq ID NO: Protein Acc MQAEGQGTGG PRGPHCGAAS FLPVFLAQAP Seq ID NO: Nucleic Ac: Coding seq ACAGCGGAGC AGACAGAGACT ATGGGAAGTC ATGGGAAGTC GCTTTTACGG CCAGATGCGA ACCAGAGACT ACGCGGGCCC CAGGTTACTA	CCTGACCTTC GCACAGGGGG GCACAGGGGGC TGCTCAGCTTC TGTCTCCAGCA CTCAGCTC TTGTCGCTG 589 Protei cession #: I 11 STGDADGPGG AQDGRCPCGA SGQRR 590 DNA se id Accession uence: 90 11 GCAGAGTGAG TCAGCGCC CCTCCTCCCCC CCTCCTGCCC CCTCCTGCCC CCTCCTGCCC CCTCCTGCCC CCTCCTGCCC CCAGGCAGTGT CAACTGCAAT GCAGAGGAAC CCGATGTCTG	TCTCTGAGAG TCTCTGAGAG TCTGACGGCC AGCAAGGGCC TGCGCAGGAT CCGACTGACT CCTCAGGCAG AGGGAATGGT AGGAGACGG in sequence be sequence polypoperation accaccac aguence #: NM_0055 accaccacc CGCACCGCA AGCACCAAC GGCACCGCA AGCACCACC AACTTTGATC AACCACCAAC CAAGGCCCCA AGGGACCGCT AAGTGTGACT AAGCCACCTG AGGGACCACT CCAGGCTTCC AAGTGTGACT AGGCACCTC CGGGGGAACC CCAGGCTTCC CGGGGGGAACC CCGGGGGAACC CCGGGGGAACC CCGGGGGAACC CCGGGGGAACC CCGGGGGAACC	CCGGGCAGAG GATGCTGATG CCAGGAGAGG GCAGGAGAGG GCAGGAGCC TTGATGTGGA AGGCGCTAAG CCCAGCACGA 31 AGGPGEAGAT RLTAADHRQL 31 CCAGGGGCCG TGCCTGCGCT TCACTCCAG GGGAACTTCA ATGGCATCA GTTTGCCCTG GCTGCCT CCACCTCCAG GTGACCCTGC TCACTCCAG TTACATGTA ATGGCATTCA ATGGCATCA TTGCCCTG TTACTGAGA CTTACCAGG	GCTCCGGAGC GCCCAGGAGG CGGGTGCCAC GAGGAGGGCGC CCTGCGGGGC CCTGCGGGGC CCAGCTT TCACGCAGTT TCACGCAGTT TGTTTCTGTA 41	CATGCAGGCC CCCTGGCATT GGGCGGCATA GGGCGGCAGA CCCGCGGGGT CAGGAGGCCG CATTCTGCCC CGCCCTTCC CATTGTGGG GAAAATAAAG 51 ARASGPRGGA LSLLMWITQC 51 CCTGCAGCGG TGCTGCCTCT TGTGATTGCA GGTAATGGAT TGCAAGAGT TCCAAGAGT TCCAGCAGAG GGCCCTATG AGGGTTCGAT TTCTGCTATG	120 180 240 360 420 480 540 600 120 180 240 360

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Seq ID NO: 611 Protein sequence Protein Accession #: BAB84587.1

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WO 02/086443

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       DCPDGSDEEN CTANPLLCST ARYHCKNGLC IDKSPICDGQ NNCQDNSDEE SCESSQEPGS
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       GOVFVTSENQ LVYYPSITYA IIGSSVIFVL VVALLALVLH HQRKRNNLMT LPVHRLQHPV
                                                                              180
       LLSRLVVLDH PHHCNVTYNV NNGIQYVASQ AEQNASEVGS PPSYSEALLD QRPAWYDLPP
                                                                              240
       PPYSSDTESL NQADLPPYRS RSGSANSASS QAASSLLSVE DTSHSPGQPG PQEGTAEPRD
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        GCAGCCATTG CTGACTCGTG GTTGGGAAAA TTCAAGACAA TCATCTATCT CTCCTTGGTG
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        TATGTGCTTG GCCATGTGAT CAAGTCCTTG GGTGCCTTAC CAATACTGGG AGGACAAGTG
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                                                                               540
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        ACTAGATACT TCTCAGTCTT CTACCTGTCC ATCAATGCAG GGAGCTTGAT TTCTACATTT
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TTTGGAGTTC CAGGACTGCT CATGGTAATT GCACTTGTTG TGTTTGCAAT GGGAAGCAAA
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                                                                               780
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                                                                               840
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        CTAGACTGGG CAGCTGAGAA ATATCCAAAG CAGCTCATTA TGGATGTAAA GGCACTGACC AGGGTACTAT TCCTTTATAT CCCATTGCC ATGTTCTGGG CTCTTTTGGA TCAGCAGGGT
                                                                               900
                                                                               960
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         CCGGACCAGA TGCAGGTTCT AAATCCCTTT CTGGTTCTTA TCTTCATCCC GTTGTTTGAC
        TTTGTCATTT ATCGTCTGGT CTCCAAGTGT GGAATTAACT TCTCATCACT TAGGAAAATG
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        GCTGTTGGTA TGATCCTAGC GTGCCTGGCA TTTGCAGTTG CGGCAGCTGT AGAGATAAAA
                                                                              1200
         ATAAATGAAA TGGCCCCAGC CCAGTCAGGT CCCCAGGAGG TTTTCCTACA AGTCTTGAAT
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         CTGGCAGATG ATGAGGTGAA GGTGACAGTG GTGGGAAATG AAAACAATTC TCTGTTGATA
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         GAGTCCATCA AATCCTTTCA GAAAACACCA CACTATTCCA AACTGCACCT GAAAACAAAA
                                                                              1380
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                                                                              1440
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         GTGCAGGAGA AGAACTGGTA CAGTCTTGTC ATTCGTGAAG ATGGGAACAG TATCTCCAGC
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         ATGATGGTAA AGGATACAGA AAGCAAAACA ACCAATGGGA TGACAACCGT GAGGTTTGTT
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         AACACTTTGC ATAAAGATGT CAACATCTCC CTGAGTACAG ATACCTCTCT CAATGTTGGT
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         TGTAGAACAG AAGATAAGAA CTTTTCTCTG AATTTGGGTC TTCTAGACTT TGGTGCAGCA
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         TATCTGTTTG TTATTACTAA TAACACCAAT CAGGGTCTTC AGGCCTGGAA GATTGAAGAC
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         ATTCCAGCCA ACAAAATGTC CATTGCGTGG CAGCTACCAC AATATGCCCT GGTTACAGCT
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         GGGGAGGTCA TGTTCTCTGT CACAGGTCTT GAGTTTTCTT ATTCTCAGGC TCCCTCTAGC
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         CTTGTTGTGG CACAGTTCAG TGGCCTGGTA CAGTGGGCCG AATTCATTTT GTTTTCCTGC
                                                                              2040
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         CTCCTGCTGG TGATCTGCCT GATCTTCTCC ATCATGGGCT ACTACTATGT TCCTGTAAAG
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         AAACTAGAGA CCAAGAAGAC AAAACTCTGA
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         Seq ID NO: 681 Protein sequence
         Protein Accession #: AAB34388.1
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WO 02/086443
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       YVLGHVIKSL GALPILGGQV VHTVLSLIGL SLIALGTGGI KPCVAAFGGD QFEEKHAEER
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       TRYFSVFYLS INAGSLISTF ITPMLRGDVQ CFGEDCYALA FGVPGLLMVI ALVVPAMGSK
                                                                            240
       IYNKPPPEGN IVAQVPKCIW FAISNRFKNR SGDIPKRQHW LDWAAEKYPK QLIMDVKALT
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       RVLFLYIPLP MFWALLDQQG SRWTLQAIRM NRNLGFFVLQ PDQMQVLNPF LVLIFIPLFD
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       FVIYRLVSKC GINFSSLRKM AVGMILACLA FAVAAAVEIK INEMAPAQSG PQEVFLQVLN
                                                                            420
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       LADDEVKYTY VGNENNSLLI ESIKSFQKTP HYSKLHLKTK SQDFHFHLKY HNLSLYTEHS
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       VQEKNWYSLV IREDGNSISS MMVKDTESKT TNGMTTVRFV NTLHKDVNIS LSTDTSLNVG
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       EDYGVSAYRT VQRGEYPAVH CRTEDKNFSL NLGLLDFGAA YLFVITNNTN QGLQAWKIED
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       IPANKMSIAW QLPQYALVTA GEVMFSVTGL EFSYSQAPSS MKSVLQAAWL LTIAVGNIIV
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       LVVAQFSGLV QWAEFILFSC LLLVICLIFS IMGYYYVPVK TEDMRGPADK HIPHIQGNMI
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                                                                           180
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       CTTGGGAGAC AGCGGGGAGT ACAAGATGAT TCTTGTGGTT CGAAATGACT TAAAGATGGG
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       CAAAGCTCCT GATGAAGAAA CCCTGATTGC ATTATTGGCC CATGCAAAAA TGCTGGGACT
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       TTACTAGGTG GACTTTGATA TGACAACAAC CCCTCCATCA CAAGTGTTTG AAGCCTGTCA
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       PDEETLIALL AHAKMLGLTV SLIQDAGRTQ IAPGSQTVLG IGPGPADLID KVTGHLKLY
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       GGCCGAGGCG AGCCGCGCAA GTTTCCCGGG ACCCTCAGAG TTGCACTCCG AAGACTCCAG
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       CTGGGAAGAT TCGAACACCG ACCTCGTCCC GGCCCCTGCA GTCCGGATAC TCACGCCAGA
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       AGTGCGGCTG GGATCCGGCG GCCACCTGCA CCTGCGTATC TCTCGGGCCG CCCTTCCCGA
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       CCGCAGAGCG CGTGCGCGCA ACGGGGACGA CTGTCCGCTC GGGCCCGGGC GTTGCTGCCG
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       GTCGCTCCAG ACCTATGATG ACTTGTTAGC CAAAGACTGC CACTGCATAT GAGCAGTCCT
                                                                            960
       GGTCCTTCCA CTGTGCACCT GCGCGGGGGA GGCGACCTCA GTTGTCCTGC CCTGTGGAAT
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                                                                           1080
                                                                           1140
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       AAAA
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       Seq ID NO: 685 Protein sequence
       Protein Accession #: NP_004855.1
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       EDLLTRLRAN QSWEDSNTDL VPAPAVRILT PEVRLGSGGH LHLRISRAAL PEGLPEASRL
                                                                            120
       HRALFRLSPT ASRSWDVTRP LRRQLSLARP QAPALHLRLS PPPSQSDQLL AESSSARPQL
                                                                            180
       ELHLRPQAAR GRRRARARNG DDCPLGPGRC CRLHTVRASL EDLGWADWVL SPREVQVTMC
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       IGACPSQFRA ANMHAQIKTS LHRLKPDTEP APCCVPASYN PMVLIQKTDT GVSLQTYDDL
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Seq ID NO: 686 DNA sequence

Nucleic Acid Accession #: NM_002423.2 Coding sequence: 48..851

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                                                                             120
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       TCTTTGGCCT ACCTATAACT GGAATGTTAA ACTCCCGCGT CATAGAAATA ATGCAGAAGC
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       CCAGATGTGG AGTGCCAGAT GTTGCAGAAT ACTCACTATT TCCAAATAGC CCAAAATGGA
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                                                                             480
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                                                                             540
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       ACTCCTACCC ATTTGATGGG CCAGGAAACA CGCTGGCTCA TGCCTTTGCG CCTGGGACAG
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                                                                             660
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                                                                             780
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                                                                             840
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                                                                             900
                                                                             960
       CTTTTTTATT GCAGTTGGTT TTTGAATGTC TTTCACTCCT TTTATTGGTT AAACTCCTTT
                                                                            1020
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       ATGTTACATA CACAAATAAA TAAAATGTTT ATTCCATGGT AAATTTA
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                                                    41
                                                               51
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       EMQKFFGLPI TGMLNSRVIE IMQKPRCGVP DVAEYSLPPN SPKWTSKVVT YRIVSYTRDL
                                                                             120
       PHITVDRLVS KALNMWGKEI PLHFRKVVWG TADIMIGFAR GAHGDSYPFD GPGNTLAHAF
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       APGTGLGGDA HFDEDERWTD GSSLGINFLY AATHELGHSL GMGHSSDPNA VMYPTYGNGD
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       Nucleic Acid Accession #: NM_005221.3
       Coding sequence: 1..870
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                                                                             120
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       TCAGCTACCG ATTCTGACTA CTACAGCCCT ACGGGGGGAG CCCCGCACGG CTACTGCTCT
                                                                             180
       CCTACCTCGG CTTCCTATGG CAAAGCTCTC AACCCCTACC AGTATCAGTA TCACGGCGTG
                                                                             240
       AACGGCTCCG CCGGGAGCTA CCCAGCCAAA GCTTATGCCG ACTATAGCTA CGCTAGCTCC
                                                                             300
       TACCACCAGT ACGGCGGCGC CTACAACCGC GTCCCAAGCG CCACCAACCA GCCAGAGAAA
                                                                             360
       GAAGTGACCG AGCCCGAGGT GAGAATGGTG AATGGCAAAC CAAAGAAAGT TCGTAAACCC
                                                                             420
50
       AGGACTATTT ATTCCAGCTT TCAGCTGGCC GCATTACAGA GAAGGTTTCA GAAGACTCAG
                                                                             480
       TACCTCGCCT TGCCGGAACG CGCCGAGCTG GCCGCCTCGC TGGGATTGAC ACAAACACAG
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       GTGAAAATCT GGTTTCAGAA CAAAAGATCC AAGATCAAGA AGATCATGAA AAACGGGGAG
                                                                             600
       ATGCCCCCGG AGCACAGTCC CAGCTCCAGC GACCCAATGG CGTGTAACTC GCCGCAGTCT
                                                                             660
       CCAGCGGTGT GGGAGCCCCA GGGCTCGTCC CGCTCGCTCA GCCACCACCC TCATGCCCAC
                                                                             720
55
       CCTCCGACCT CCAACCAGTC CCCAGCGTCC AGCTACCTGG AGAACTCTGC ATCCTGGTAC
       ACAAGTGCAG CCAGCTCAAT CAATTCCCAC CTGCCGCCGC CGGGCTCCTT ACAGCACCCG
                                                                             840.
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                                                                             120
       EVTEPEVRMV NGKPKKVRKP RTIYSSFQLA ALQRRFQKTQ YLALPERAEL AASLGLTQTQ
                                                                             180
       VKIWFQNKRS KIKKIMKNGE MPPEHSPSSS DPMACNSPQS PAVWEPQGSS RSLSHHPHAH
                                                                             240
       PPTSNOSPAS SYLENSASWY TSAASSINSH LPPPGSLOHP LALASGTLY
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It is understood that the examples described above in no way serve to limit the true scope of this invention, but rather are presented for illustrative purposes. All publications, sequences of accession numbers, and patent applications cited in this specification are herein neorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference.

WHAT IS CLAIMED IS:

1	1.	A method of detecting a lung cancer-associated transcript in a cell		
2	from a patient, the method comprising contacting a biological sample from the patient with a			
3	polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence			
4	as shown in Table	s 1A-16.		
1	2.	The method of claim 1, wherein the polynucleotide selectively		
2	hybridizes to a sec	quence at least 95% identical to a sequence as shown in Tables 1A-16.		
1	3.	The method of claim 1, wherein the biological sample is a tissue		
2	sample.			
1	4.	The method of claim 1, wherein the biological sample comprises		
2	isolated nucleic ac	eids.		
1	5.	The method of claim 4, wherein the nucleic acids are mRNA.		
1	6.	The method of claim 4, further comprising the step of amplifying		
2	nucleic acids befo	re the step of contacting the biological sample with the polynucleotide.		
1	7.	The method of claim 1, wherein the polynucleotide comprises a		
2	sequence as show	n in Tables 1A-16.		
1	8.	The method of claim 1, wherein the polynucleotide is labeled.		
1	9.	The method of claim 8, wherein the label is a fluorescent label.		
1	10	The method of claim 1, wherein the polynucleotide is immobilized on		
2	a solid surface.			
1	11	The method of claim 1, wherein the patient is undergoing a therapeutic		
2	regimen to treat lu	nng cancer.		
1	. 12	The method of claim 1, wherein the patient is suspected of having lung		
2	cancer.			
1	13	A method of monitoring the efficacy of a therapeutic treatment of lung		
2	cancer, the metho	d comprising the steps of:		

PCT/US02/12476 WO 02/086443 3 (i) providing a biological sample from a patient undergoing the therapeutic 4 treatment; and 5 (ii) determining the level of a lung cancer-associated transcript in the 6 biological sample by contacting the biological sample with a polynucleotide that selectively 7 hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, 8 thereby monitoring the efficacy of the therapy. 1 . 14. The method of claim 13, further comprising the step of: (iii) comparing 2 the level of the lung cancer-associated transcript to a level of the lung cancer-associated 3 transcript in a biological sample from the patient prior to, or earlier in, the therapeutic 4 treatment. 1 15. The method of claim 13, wherein the patient is a human. 1 16. A method of monitoring the efficacy of a therapeutic treatment of lung 2 cancer, the method comprising the steps of: 3 (i) providing a biological sample from a patient undergoing the therapeutic 4 treatment; and 5 (ii) determining the level of a lung cancer-associated antibody in the biological 6 sample by contacting the biological sample with a polypeptide encoded by a polynucleotide 7 that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in 8 Tables 1A-16, wherein the polypeptide specifically binds to the lung cancer-associated 9 antibody, thereby monitoring the efficacy of the therapy. 1 17. The method of claim 16, further comprising the step of: (iii) comparing 2 the level of the lung cancer-associated antibody to a level of the lung cancer-associated 3 antibody in a biological sample from the patient prior to, or earlier in, the therapeutic 4 treatment. 1 18. The method of claim 16, wherein the patient is a human. 1 19. A method of monitoring the efficacy of a therapeutic treatment of lung 2 cancer, the method comprising the steps of: 3

4

treatment; and

(i) providing a biological sample from a patient undergoing the therapeutic

5	(ii) determining the level of a lung cancer-associated polypeptide in the			
6	biological sample by contacting the biological sample with an antibody, wherein the antibody			
7	specifically binds to a polypeptide encoded by a polynucleotide that selectively hybridizes to			
8	a sequence at least 80% identical to a sequence as shown in Tables 1A-16, thereby			
9	monitoring the efficacy of the therapy.			
1	20.	The method of claim 19, further comprising the step of: (iii) comparing		
2	the level of the lung cancer-associated polypeptide to a level of the lung cancer-associated			
3	polypeptide in a biological sample from the patient prior to, or earlier in, the therapeutic			
4	treatment.			
1	21.	The method of claim 19, wherein the patient is a human.		
1	22.	An isolated nucleic acid molecule consisting of a polynucleotide		
2	sequence as shown in Tables 1A-16.			
	•			
1	23.	The nucleic acid molecule of claim 22, which is labeled.		
1	24.	The nucleic acid of claim 23, wherein the label is a fluorescent label		
1	25.	An expression vector comprising the nucleic acid of claim 22.		
1	26.	A host cell comprising the expression vector of claim 25.		
1	27.	An isolated polypeptide which is encoded by a nucleic acid molecule		
2	having polynucleotic	le sequence as shown in Tables 1A-16.		
1	28.	An antibody that specifically binds a polypeptide of claim 27.		
1	29.	The antibody of claim 28, further conjugated to an effector component.		
1	30.	The antibody of claim 29, wherein the effector component is a		
2	fluorescent label.			
	21	The autiliardy of alaim 20 whomain the affector component is a		
1	31.	The antibody of claim 29, wherein the effector component is a		
2	radioisotope or a cyt	otoxic chemical.		
1	. 32	The antibody of claim 29, which is an antibody fragment.		

1	33. The antibody of claim 29, which is a numarized antibody				
1	34. A method of detecting a lung cancer cell in a biological sample fro	m a			
2	patient, the method comprising contacting the biological sample with an antibody of clair	n			
3	28.				
1	35. The method of claim 34, wherein the antibody is further conjugate	d to			
2	an effector component.				
1	36. The method of claim 35, wherein the effector component is a				
2	fluorescent label.				
1	37. A method of detecting antibodies specific to lung cancer in a patient	nt,			
2	the method comprising contacting a biological sample from the patient with a polypeptide	€			
3	encoded by a nucleic acid comprises a sequence from Tables 1A-16.				
1	38. A method for identifying a compound that modulates a lung cancer	r-			
2	associated polypeptide, the method comprising the steps of:				
3	(i) contacting the compound with a lung cancer-associated polypeptide, th	е			
4	polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least				
5	80% identical to a sequence as shown in Tables 1A-16; and				
6	(ii) determining the functional effect of the compound upon the polypeptic	le.			
1	39. The method of claim 38, wherein the functional effect is a physical	l			
2	effect.				
1	40. The method of claim 38, wherein the functional effect is a chemical	ıl			
2	effect.				
1	41. The method of claim 38, wherein the polypeptide is expressed in a				
2	eukaryotic host cell or cell membrane.				
1	42. The method of claim 38, wherein the functional effect is determine	d by			
2	measuring ligand binding to the polypeptide.				
1	The method of claim 38 wherein the polymentide is recombinant				

l	44.	A method of inhibiting proliferation of a fully calicel-associated cell to	
2	treat lung cancer in a	a patient, the method comprising the step of administering to the subject a	
3	therapeutically effective amount of a compound identified using the method of claim 38.		
1	45.	The method of claim 44, wherein the compound is an antibody.	
1	46.	The method of claim 45, wherein the patient is a human.	
1	47.	A drug screening assay comprising the steps of	
2	(i) ad	ministering a test compound to a mammal having lung cancer or a cell	
3	isolated therefrom;		
4	(ii) comparing the level of gene expression of a polynucleotide that selectively		
5	hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16 in a		
6	treated cell or mammal with the level of gene expression of the polynucleotide in a control		
7	cell or mammal, wherein a test compound that modulates the level of expression of the		
8	polynucleotide is a c	andidate for the treatment of lung cancer.	
1	48.	The assay of claim 47, wherein the control is a mammal with lung	
2	cancer or a cell there	efrom that has not been treated with the test compound.	
1	49.	The assay of claim 47, wherein the control is a normal cell or mammal.	
1	50.	A method for treating a mammal having lung cancer comprising	
2	administering a com	pound identified by the assay of claim 47.	
1	51.	A pharmaceutiPcal composition for treating a mammal having lung	
2	cancer, the composition comprising a compound identified by the assay of claim 47 and a		
3	physiologically acceptable excipient.		